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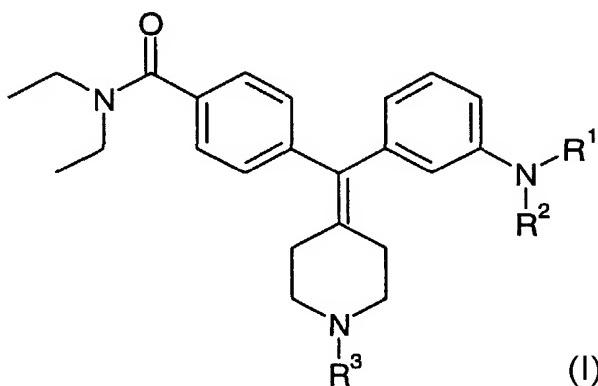
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(54) Title: PHENYL-PIPERIDIN-4-YLIDENE-METHYL-BENZAMIDE DERIVATIVES FOR THE TREATMENT OF PAIN OR GASTROINTESTINAL DISORDERS

(57) Abstract: Compounds of general formula: wherein R¹, R² and R³ are as defined in the specification, as well as salts, enantiomers thereof and pharmaceutical compositions including the compounds are prepared. They are useful in therapy, in particular in the management of pain.



WO 2004/041784 A1

PHENYL-PIPERIDIN-4-YLIDENE-METHYL-BENZAMIDE DERIVATIVES FOR THE
TREATMENT OF PAIN OR GASTROINTESTINAL DISORDERS

BACKGROUND OF THE INVENTION

1. Field of the Invention

5 The present invention is directed to novel compounds, to a process for their preparation, their use and pharmaceutical compositions comprising the novel compounds. The novel compounds are useful in therapy, and in particular for the treatment of pain, anxiety and functional gastrointestinal disorders.

2. Discussion of Relevant Art

10 The δ receptor has been identified as having a role in many bodily functions such as circulatory and pain systems. Ligands for the δ receptor may therefore find potential use as analgesics, and/or as antihypertensive agents. Ligands for the δ receptor have also been shown to possess immunomodulatory activities.

15 The identification of at least three different populations of opioid receptors (μ , δ and κ) is now well established and all three are apparent in both central and peripheral nervous systems of many species including man. Analgesia has been observed in various animal models when one or more of these receptors has been activated.

20 With few exceptions, currently available selective opioid δ ligands are peptidic in nature and are unsuitable for administration by systemic routes. One example of a non-peptidic δ -agonist is SNC80 (Bilsky E.J. et al., Journal of Pharmacology and Experimental Therapeutics, 273(1), pp. 359-366 (1995)).

25 Many δ agonist compounds that have been identified in the prior art have many disadvantages in that they suffer from poor pharmacokinetics and are not analgesic when administered by systemic routes. Also, it has been documented that many of these δ agonist compounds show significant convulsive effects when administered systemically.

U.S. Patent No. 6,187,792 to Delorme et al. describes some δ -agonists.

However, there is still a need for improved δ -agonists.

DETAILED DESCRIPTION OF THE INVENTION**Definitions**

Unless specified otherwise within this specification, the nomenclature used in this specification generally follows the examples and rules stated in *Nomenclature of Organic Chemistry, Sections A, B, C, D, E, F, and H*, Pergamon Press, Oxford, 1979, which is incorporated by references herein for its exemplary chemical structure names and rules on naming chemical structures.

The term "C_{m-n}" or "C_{m-n} group" used alone or as a prefix, refers to any group having m to n carbon atoms.

10 The term "hydrocarbon" used alone or as a suffix or prefix, refers to any structure comprising only carbon and hydrogen atoms up to 14 carbon atoms.

The term "hydrocarbon radical" or "hydrocarbyl" used alone or as a suffix or prefix, refers to any structure as a result of removing one or more hydrogens from a hydrocarbon.

15 The term "alkyl" used alone or as a suffix or prefix, refers to monovalent straight or branched chain hydrocarbon radicals comprising 1 to about 12 carbon atoms. An "alkyl" may optionally contain one or more unsaturated carbon-carbon bonds.

20 The term "alkylene" used alone or as suffix or prefix, refers to divalent straight or branched chain hydrocarbon radicals comprising 1 to about 12 carbon atoms, which serves to links two structures together.

The term "alkenyl" used alone or as suffix or prefix, refers to a monovalent straight or branched chain hydrocarbon radical having at least one carbon-carbon double bond and comprising at least 2 up to about 12 carbon atoms.

25 The term "alkynyl" used alone or as suffix or prefix, refers to a monovalent straight or branched chain hydrocarbon radical having at least one carbon-carbon triple bond and comprising at least 2 up to about 12 carbon atoms.

The term "cycloalkyl," used alone or as suffix or prefix, refers to a monovalent ring-containing hydrocarbon radical comprising at least 3 up to about 12 carbon atoms.

The term "cycloalkenyl" used alone or as suffix or prefix, refers to a monovalent ring-containing hydrocarbon radical having at least one carbon-carbon double bond and comprising at least 3 up to about 12 carbon atoms.

5 The term "cycloalkynyl" used alone or as suffix or prefix, refers to a monovalent ring-containing hydrocarbon radical having at least one carbon-carbon triple bond and comprising about 7 up to about 12 carbon atoms.

10 The term "aryl" used alone or as suffix or prefix, refers to a monovalent hydrocarbon radical having one or more polyunsaturated carbon rings having aromatic character, (e.g., $4n + 2$ delocalized electrons) and comprising 5 up to about 14 carbon atoms.

The term "arylene" used alone or as suffix or prefix, refers to a divalent hydrocarbon radical having one or more polyunsaturated carbon rings having aromatic character, (e.g., $4n + 2$ delocalized electrons) and comprising 5 up to about 14 carbon atoms, which serves to link two structures together.

15 The term "heterocycle" used alone or as a suffix or prefix, refers to a ring-containing structure or molecule having one or more multivalent heteroatoms, independently selected from N, O, P and S, as a part of the ring structure and including at least 3 and up to about 20 atoms in the ring(s). Heterocycle may be saturated or unsaturated, containing one or more double bonds, and heterocycle may 20 contain more than one ring. When a heterocycle contains more than one ring, the rings may be fused or unfused. Fused rings generally refer to at least two rings share two atoms therebetween. Heterocycle may have aromatic character or may not have aromatic character.

25 The term "heteroaromatic" used alone or as a suffix or prefix, refers to a ring-containing structure or molecule having one or more multivalent heteroatoms, independently selected from N, O, P and S, as a part of the ring structure and including at least 3 and up to about 20 atoms in the ring(s), wherein the ring-containing structure or molecule has an aromatic character (e.g., $4n + 2$ delocalized electrons).

30 The term "heterocyclic group," "heterocyclic moiety," "heterocyclic," or "heterocyclo" used alone or as a suffix or prefix, refers to a radical derived from a heterocycle by removing one or more hydrogens therefrom.

The term "heterocyclyl" used alone or as a suffix or prefix, refers a monovalent radical derived from a heterocycle by removing one hydrogen therefrom.

The term "heterocyclylene" used alone or as a suffix or prefix, refers to a divalent radical derived from a heterocycle by removing two hydrogens therefrom, which serves to links two structures together.

The term "heteroaryl" used alone or as a suffix or prefix, refers to a heterocyclyl having aromatic character.

The term "heterocylcoalkyl" used alone or as a suffix or prefix, refers to a heterocyclyl that does not have aromatic character.

The term "heteroarylene" used alone or as a suffix or prefix, refers to a heterocyclylene having aromatic character.

The term "heterocycloalkylene" used alone or as a suffix or prefix, refers to a heterocyclylene that does not have aromatic character.

The term "six-membered" used as prefix refers to a group having a ring that contains six ring atoms.

The term "five-membered" used as prefix refers to a group having a ring that contains five ring atoms.

A five-membered ring heteroaryl is a heteroaryl with a ring having five ring atoms wherein 1, 2 or 3 ring atoms are independently selected from N, O and S.

Exemplary five-membered ring heteroaryls are thienyl, furyl, pyrrolyl, imidazolyl, thiazolyl, oxazolyl, pyrazolyl, isothiazolyl, isoxazolyl, 1,2,3-triazolyl, tetrazolyl, 1,2,3-thiadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-triazolyl, 1,2,4-thiadiazolyl, 1,2,4-oxadiazolyl, 1,3,4-triazolyl, 1,3,4-thiadiazolyl, and 1,3,4- oxadiazolyl.

A six-membered ring heteroaryl is a heteroaryl with a ring having six ring atoms wherein 1, 2 or 3 ring atoms are independently selected from N, O and S.

Exemplary six-membered ring heteroaryls are pyridyl, pyrazinyl, pyrimidinyl, triazinyl and pyridazinyl.

The term "substituted" used as a prefix refers to a structure, molecule or group, wherein one or more hydrogens are replaced with one or more C₁₋₆hydrocarbon groups, or one or more chemical groups containing one or more heteroatoms selected from N, O, S, F, Cl, Br, I, and P. Exemplary chemical groups containing one or more heteroatoms include -NO₂, -OR, -Cl, -Br, -I, -F, -CF₃,

-C(=O)R, -C(=O)OH, -NH₂, -SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR₂, -NRC(=O)R, oxo (=O), imino (=NR), thio (=S), and oximino (=N-OR), wherein each "R" is a C₁₋₆hydrocarbyl. For example, substituted phenyl may refer to nitrophenyl, methoxyphenyl, chlorophenyl, aminophenyl, etc., 5 wherein the nitro, methoxy, chloro, and amino groups may replace any suitable hydrogen on the phenyl ring.

The term "substituted" used as a suffix of a first structure, molecule or group, followed by one or more names of chemical groups refers to a second structure, molecule or group, which is a result of replacing one or more hydrogens of the first 10 structure, molecule or group with the one or more named chemical groups. For example, a "phenyl substituted by nitro" refers to nitrophenyl.

Heterocycle includes, for example, monocyclic heterocycles such as: aziridine, oxirane, thiirane, azetidine, oxetane, thietane, pyrrolidine, pyrroline, imidazolidine, pyrazolidine, pyrazoline, dioxolane, sulfolane 2,3-dihydrofuran, 2,5-dihydrofuran tetrahydrofuran, thiophane, piperidine, 1,2,3,6-tetrahydro-pyridine, 15 piperazine, morpholine, thiomorpholine, pyran, thiopyran, 2,3-dihydropyran, tetrahydropyran, 1,4-dihydropyridine, 1,4-dioxane, 1,3-dioxane, dioxane, homopiperidine, 2,3,4,7-tetrahydro-1*H*-azepine homopiperazine, 1,3-dioxepane, 4,7-dihydro-1,3-dioxepin, and hexamethylene oxide.

In addition, heterocycle includes aromatic heterocycles, for example, pyridine, 20 pyrazine, pyrimidine, pyridazine, thiophene, furan, furazan, pyrrole, imidazole, thiazole, oxazole, pyrazole, isothiazole, isoxazole, 1,2,3-triazole, tetrazole, 1,2,3-thiadiazole, 1,2,3-oxadiazole, 1,2,4-triazole, 1,2,4-thiadiazole, 1,2,4-oxadiazole, 1,3,4-triazole, 1,3,4-thiadiazole, and 1,3,4- oxadiazole.

Additionally, heterocycle encompass polycyclic heterocycles, for example, 25 indole, indoline, isoindoline, quinoline, tetrahydroquinoline, isoquinoline, tetrahydroisoquinoline, 1,4-benzodioxan, coumarin, dihydrocoumarin, benzofuran, 2,3-dihydrobenzofuran, isobenzofuran, chromene, chroman, isochroman, xanthene, phenoxathiin, thianthrene, indolizine, isoindole, indazole, purine, phthalazine, 30 naphthyridine, quinoxaline, quinazoline, cinnoline, pteridine, phenanthridine, perimidine, phenanthroline, phenazine, phenothiazine, phenoxazine, 1,2-benzisoxazole, benzothiophene, benzoxazole, benzthiazole, benzimidazole,

6

benztriazole, thioxanthine, carbazole, carboline, acridine, pyrolizidine, and quinolizidine.

In addition to the polycyclic heterocycles described above, heterocycle includes polycyclic heterocycles wherein the ring fusion between two or more rings includes more than one bond common to both rings and more than two atoms common to both rings. Examples of such bridged heterocycles include quinuclidine, diazabicyclo[2.2.1]heptane and 7-oxabicyclo[2.2.1]heptane.

Heterocyclyl includes, for example, monocyclic heterocyclyls, such as: aziridinyl, oxiranyl, thiiranyl, azetidinyl, oxetanyl, thietanyl, pyrrolidinyl, pyrrolinyl, 10 imidazolidinyl, pyrazolidinyl, pyrazolinyl, dioxolanyl, sulfolanyl, 2,3-dihydrofuranyl, 2,5-dihydrofuranyl, tetrahydrofuranyl, thiophanyl, piperidinyl, 1,2,3,6-tetrahydro- pyridinyl, piperazinyl, morpholinyl, thiomorpholinyl, pyranyl, thiopyranyl, 2,3- dihydropyranyl, tetrahydropyranyl, 1,4-dihydropyridinyl, 1,4-dioxanyl, 1,3-dioxanyl, dioxanyl, homopiperidinyl, 2,3,4,7-tetrahydro-1*H*-azepinyl, homopiperazinyl, 1,3- 15 dioxepanyl, 4,7-dihydro-1,3-dioxepinyl, and hexamethylene oxidyl.

In addition, heterocyclyl includes aromatic heterocyclyls or heteroaryl, for example, pyridinyl, pyrazinyl, pyrimidinyl, pyridazinyl, thienyl, furyl, furazanyl, 20 pyrrolyl, imidazolyl, thiazolyl, oxazolyl, pyrazolyl, isothiazolyl, isoxazolyl, 1,2,3-triazolyl, tetrazolyl, 1,2,3-thiadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-triazolyl, 1,2,4-thiadiazolyl, 1,2,4-oxadiazolyl, 1,3,4-triazolyl, 1,3,4-thiadiazolyl, and 1,3,4-oxadiazolyl.

Additionally, heterocyclyl encompasses polycyclic heterocyclyls (including both aromatic or non-aromatic), for example, indolyl, indolinyl, isoindolinyl, 25 quinolinyl, tetrahydroquinolinyl, isoquinolinyl, tetrahydroisoquinolinyl, 1,4-benzodioxanyl, coumarinyl, dihydrocoumarinyl, benzofuranyl, 2,3-dihydrobenzofuranyl, isobenzofuranyl, chromenyl, chromanyl, isochromanyl, xanthenyl, phenoxathiinyl, thianthrenyl, indolizinyl, isoindolyl, indazolyl, purinyl, phthalazinyl, naphthyridinyl, quinoxalinyl, quinazolinyl, cinnolinyl, pteridinyl, phenanthridinyl, perimidinyl, phenanthrolinyl, phenazinyl, phenothiazinyl, 30 phenoxazinyl, 1,2-benzisoxazolyl, benzothiophenyl, benzoxazolyl, benzthiazolyl, benzimidazolyl, benztriazolyl, thioxanthinyl, carbazolyl, carbolinyl, acridinyl, pyrolizidinyl, and quinolizidinyl.

In addition to the polycyclic heterocycls described above, heterocycl includes polycyclic heterocycls wherein the ring fusion between two or more rings includes more than one bond common to both rings and more than two atoms common to both rings. Examples of such bridged heterocycles include quinuclidinyl, 5 diazabicyclo[2.2.1]heptyl; and 7-oxabicyclo[2.2.1]heptyl.

The term "alkoxy" used alone or as a suffix or prefix, refers to radicals of the general formula -O-R, wherein R is selected from a hydrocarbon radical. Exemplary alkoxy includes methoxy, ethoxy, propoxy, isopropoxy, butoxy, t-butoxy, isobutoxy, cyclopropylmethoxy, allyloxy, and propargyloxy.

10 The term "amine" or "amino" used alone or as a suffix or prefix, refers to radicals of the general formula -NRR', wherein R and R' are independently selected from hydrogen or a hydrocarbon radical.

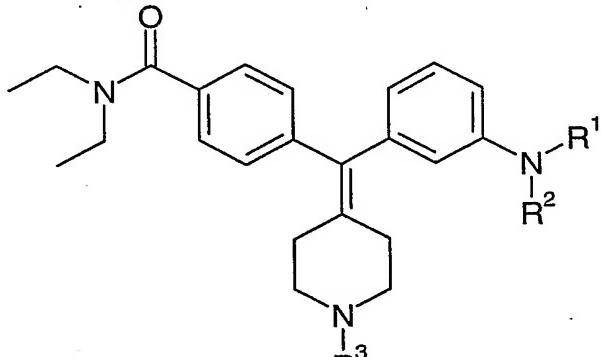
Halogen includes fluorine, chlorine, bromine and iodine.

15 "Halogenated," used as a prefix of a group, means one or more hydrogens on the group is replaced with one or more halogens.

"RT" or "rt" means room temperature.

Description of Embodiments

In one aspect, the invention provides a compound of formula I, diastereomers 20 thereof and pharmaceutically acceptable salts thereof:



I

wherein

R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, $R^8-C(=O)-$, $R^8-S(=O)_2-$, $R^8-S(=O)-$, $R^8-NHC(=O)-$, $R^8-C(=S)-$ and $R^8-NH-C(=S)-$, wherein R^8 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl-
5 C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-9} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-9} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl used in defining R^1 and R^8 are optionally substituted with one or more groups selected from -R, -NO₂, -OR, -Cl, -Br, -I, -F, -CF₃, -C(=O)R, -C(=O)OH, -NH₂, -SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(=O)R, -CN,
10 -OH, -C(=O)OR, -C(=O)NR₂, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, selected from -H, C_{1-6} alkyl and phenyl;

R^2 is selected from -H and C_{1-6} alkyl optionally substituted with one or more groups selected from halogen, -CF₃, -OH, C_{1-3} alkoxy, and halogen; and

R^3 is selected from -H, C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, -NO₂, -CF₃, C_{1-6} alkoxy and halogen.

In one embodiment, the compounds of the present invention are represented by formula I, wherein

R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-4} alkyl, halogen, -CF₃, -OH, C_{1-3} alkoxy, phenoxy, and
25 halogen;

R^2 is selected from -H and C_{1-3} alkyl; and

R^3 is selected from -H and C_{1-6} alkyl-O-C(=O)-.

In another embodiment, the compounds of the present invention are represented by formula I, wherein R^1 is R^9-CH_2- , wherein R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted
30

with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy and halogen; and

R² and R³ are hydrogen.

In another embodiment, the compounds of the present invention are
5 represented by formula I,

wherein R¹ is R⁹-CH₂-, wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl, optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

10 R² and R³ are hydrogen.

In another embodiment, the compounds of the present invention are
represented by formula I, wherein

wherein R¹ is R⁹-CH₂-, wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl; and

15 R² and R³ are hydrogen.

In another embodiment, the compounds of the present invention are
represented by formula I, wherein

R¹ is selected from C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl,
wherein said C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally
20 substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

R² is -H or C₁₋₃alkyl; and

25 R³ is -H, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said
C₁₋₆alkyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one
or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and
halogen.

In another embodiment, the compounds of the present invention are
represented by formula I, wherein

30 R¹ is selected from 1-propyl, 2-propyl, 1-butyl, 2-butyl, t-butyl, 2-methyl-1-
propyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, and cyclononyl;

R² is selected from -H, methyl, ethyl, 1-propyl and 2-propyl; and

10

R^3 is selected from -H, methyl, ethyl, allyl, 3,3-dimethyl-allyl, cyclopropylmethyl, 2-methoxy-ethyl, and 3-methoxy-1-propyl.

In another embodiment, the compounds of the present invention are represented by formula I, wherein

- 5 R^1 is selected from $R^8-C(=O)-$, $R^8-S(=O)_2-$, $R^8-S(=O)-$, $R^8-NHC(=O)-$, R^8-
 $C(=S)-$ and $R^8-NH-C(=S)-$, wherein R^8 is selected from C_{3-6} alkyl, C_{6-10} aryl,
 C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl; wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl are
10 optionally substituted with C_{1-4} alkyl, halogen, - CF_3 , -OH, C_{1-3} alkoxy, phenoxy, and
halogen;

R^2 is -H; and

R^3 is selected from -H and C_{1-6} alkyl-O-C(=O)-.

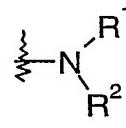
- In a further embodiment, the compounds of the present invention are
15 represented by formula I, wherein

- 15 R^1 is selected from $R^8-C(=O)-$, $R^8-S(=O)_2-$, $R^8-S(=O)-$, $R^8-NHC(=O)-$, R^8-
 $C(=S)-$ and $R^8-NH-C(=S)-$, wherein R^8 is selected from phenyl, benzyl, phenethyl and
cyclohexyl, wherein said phenyl, benzyl, phenethyl and cyclohexyl are optionally
substituted with one or more groups selected from methyl, methoxy and halogen;

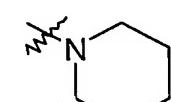
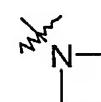
- 20 R^2 is -H; and

R^3 is selected from -H and C_{1-6} alkyl-O-C(=O)-.

In an even further embodiment, the compounds of the present invention are those of formula I, wherein



of formula I is selected from



and ; and

25

R^3 is selected from -H and C_{1-6} alkyl-O-C(=O)-.

11

It will be understood that when compounds of the present invention contain one or more chiral centers, the compounds of the invention may exist in, and be isolated as, enantiomeric or diastereomeric forms, or as a racemic mixture. The present invention includes any possible enantiomers, diastereomers, racemates or mixtures thereof, of a compound of Formula I. The optically active forms of the compound of the invention may be prepared, for example, by chiral chromatographic separation of a racemate, by synthesis from optically active starting materials or by asymmetric synthesis based on the procedures described thereafter.

It will also be appreciated that certain compounds of the present invention may exist as geometrical isomers, for example E and Z isomers of alkenes. The present invention includes any geometrical isomer of a compound of Formula I. It will further be understood that the present invention encompasses tautomers of the compounds of the formula I.

It will also be understood that certain compounds of the present invention may exist in solvated, for example hydrated, as well as unsolvated forms. It will further be understood that the present invention encompasses all such solvated forms of the compounds of the formula I.

Within the scope of the invention are also salts of the compounds of the formula I. Generally, pharmaceutically acceptable salts of compounds of the present invention may be obtained using standard procedures well known in the art, for example by reacting a sufficiently basic compound, for example an alkyl amine with a suitable acid, for example, HCl or acetic acid, to afford a physiologically acceptable anion. It may also be possible to make a corresponding alkali metal (such as sodium, potassium, or lithium) or an alkaline earth metal (such as a calcium) salt by treating a compound of the present invention having a suitably acidic proton, such as a carboxylic acid or a phenol with one equivalent of an alkali metal or alkaline earth metal hydroxide or alkoxide (such as the ethoxide or methoxide), or a suitably basic organic amine (such as choline or meglumine) in an aqueous medium, followed by conventional purification techniques.

In one embodiment, the compound of formula I above may be converted to a pharmaceutically acceptable salt or solvate thereof, particularly, an acid addition salt

12

such as a hydrochloride, hydrobromide, phosphate, acetate, fumarate, maleate, tartrate, citrate, methanesulphonate or *p*-toluenesulphonate.

The novel compounds of the present invention are useful in therapy, especially for the treatment of various pain conditions such as chronic pain, neuropathic pain, acute pain, cancer pain, pain caused by rheumatoid arthritis, migraine, visceral pain etc. This list should however not be interpreted as exhaustive.

Compounds of the invention are useful as immunomodulators, especially for autoimmune diseases, such as arthritis, for skin grafts, organ transplants and similar surgical needs, for collagen diseases, various allergies, for use as anti-tumour agents and anti viral agents.

Compounds of the invention are useful in disease states where degeneration or dysfunction of opioid receptors is present or implicated in that paradigm. This may involve the use of isotopically labelled versions of the compounds of the invention in diagnostic techniques and imaging applications such as positron emission tomography (PET).

Compounds of the invention are useful for the treatment of diarrhoea, depression, anxiety and stress-related disorders such as post-traumatic stress disorders, panic disorder, generalized anxiety disorder, social phobia, and obsessive compulsive disorder, urinary incontinence, premature ejaculation, various mental illnesses, cough, lung oedema, various gastro-intestinal disorders, e.g. constipation, functional gastrointestinal disorders such as Irritable Bowel Syndrome and Functional Dyspepsia, Parkinson's disease and other motor disorders, traumatic brain injury, stroke, cardioprotection following myocardial infarction, spinal injury and drug addiction, including the treatment of alcohol, nicotine, opioid and other drug abuse and for disorders of the sympathetic nervous system for example hypertension.

Compounds of the invention are useful as an analgesic agent for use during general anaesthesia and monitored anaesthesia care. Combinations of agents with different properties are often used to achieve a balance of effects needed to maintain the anaesthetic state (e.g. amnesia, analgesia, muscle relaxation and sedation).

Included in this combination are inhaled anaesthetics, hypnotics, anxiolytics, neuromuscular blockers and opioids.

13

Also within the scope of the invention is the use of any of the compounds according to the formula I above, for the manufacture of a medicament for the treatment of any of the conditions discussed above.

5 A further aspect of the invention is a method for the treatment of a subject suffering from any of the conditions discussed above, whereby an effective amount of a compound according to the formula I above, is administered to a patient in need of such treatment.

Thus, the invention provides a compound of formula I, or pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined for use in therapy.

10 In a further aspect, the present invention provides the use of a compound of formula I, or a pharmaceutically acceptable salt or solvate thereof, as hereinbefore defined in the manufacture of a medicament for use in therapy.

In the context of the present specification, the term "therapy" also includes "prophylaxis" unless there are specific indications to the contrary. The term 15 "therapeutic" and "therapeutically" should be construed accordingly. The term "therapy" within the context of the present invention further encompasses to administer an effective amount of a compound of the present invention, to mitigate either a pre-existing disease state, acute or chronic, or a recurring condition. This definition also encompasses prophylactic therapies for prevention of recurring 20 conditions and continued therapy for chronic disorders.

The compounds of the present invention are useful in therapy, especially for the therapy of various pain conditions including, but not limited to: chronic pain, neuropathic pain, acute pain, back pain, cancer pain, and visceral pain.

25 In use for therapy in a warm-blooded animal such as a human, the compound of the invention may be administered in the form of a conventional pharmaceutical composition by any route including orally, intramuscularly, subcutaneously, topically, intranasally, intraperitoneally, intrathoracically, intravenously, epidurally, intrathecally, intracerebroventricularly and by injection into the joints.

30 In one embodiment of the invention, the route of administration may be orally, intravenously or intramuscularly.

The dosage will depend on the route of administration, the severity of the disease, age and weight of the patient and other factors normally considered by the

attending physician, when determining the individual regimen and dosage level at the most appropriate for a particular patient.

For preparing pharmaceutical compositions from the compounds of this invention, inert, pharmaceutically acceptable carriers can be either solid and liquid.

- 5 Solid form preparations include powders, tablets, dispersible granules, capsules, cachets, and suppositories.

A solid carrier can be one or more substances, which may also act as diluents, flavoring agents, solubilizers, lubricants, suspending agents, binders, or table disintegrating agents; it can also be an encapsulating material.

- 10 In powders, the carrier is a finely divided solid, which is in a mixture with the finely divided compound of the invention, or the active component. In tablets, the active component is mixed with the carrier having the necessary binding properties in suitable proportions and compacted in the shape and size desired.

- 15 For preparing suppository compositions, a low-melting wax such as a mixture of fatty acid glycerides and cocoa butter is first melted and the active ingredient is dispersed therein by, for example, stirring. The molten homogeneous mixture is then poured into convenient sized moulds and allowed to cool and solidify.

- 20 Suitable carriers are magnesium carbonate, magnesium stearate, talc, lactose, sugar, pectin, dextrin, starch, tragacanth, methyl cellulose, sodium carboxymethyl cellulose, a low-melting wax, cocoa butter, and the like.

The term composition is also intended to include the formulation of the active component with encapsulating material as a carrier providing a capsule in which the active component (with or without other carriers) is surrounded by a carrier which is thus in association with it. Similarly, cachets are included.

- 25 Tablets, powders, cachets, and capsules can be used as solid dosage forms suitable for oral administration.

- Liquid form compositions include solutions, suspensions, and emulsions. For example, sterile water or water propylene glycol solutions of the active compounds may be liquid preparations suitable for parenteral administration. Liquid
30 compositions can also be formulated in solution in aqueous polyethylene glycol solution.

Aqueous solutions for oral administration can be prepared by dissolving the active component in water and adding suitable colorants, flavoring agents, stabilizers, and thickening agents as desired. Aqueous suspensions for oral use can be made by dispersing the finely divided active component in water together with a viscous material such as natural synthetic gums, resins, methyl cellulose, sodium carboxymethyl cellulose, and other suspending agents known to the pharmaceutical formulation art.

Depending on the mode of administration, the pharmaceutical composition will preferably include from 0.05% to 99%w (per cent by weight), more preferably from 0.10 to 50%w, of the compound of the invention, all percentages by weight being based on total composition.

A therapeutically effective amount for the practice of the present invention may be determined, by the use of known criteria including the age, weight and response of the individual patient, and interpreted within the context of the disease which is being treated or which is being prevented, by one of ordinary skills in the art.

Within the scope of the invention is the use of any compound of formula I as defined above for the manufacture of a medicament.

Also within the scope of the invention is the use of any compound of formula I for the manufacture of a medicament for the therapy of pain.

Additionally provided is the use of any compound according to Formula I for the manufacture of a medicament for the therapy of various pain conditions including, but not limited to: chronic pain, neuropathic pain, acute pain, back pain, cancer pain, and visceral pain.

A further aspect of the invention is a method for therapy of a subject suffering from any of the conditions discussed above, whereby an effective amount of a compound according to the formula I above, is administered to a patient in need of such therapy.

Additionally, there is provided a pharmaceutical composition comprising a compound of Formula I, or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable carrier.

Particularly, there is provided a pharmaceutical composition comprising a compound of Formula I, or a pharmaceutically acceptable salt thereof, in association

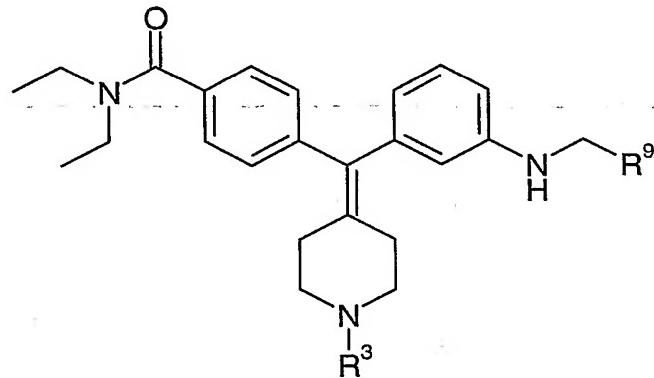
16

with a pharmaceutically acceptable carrier for therapy, more particularly for therapy of pain.

Further, there is provided a pharmaceutical composition comprising a compound of Formula I, or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable carrier use in any of the conditions discussed above.

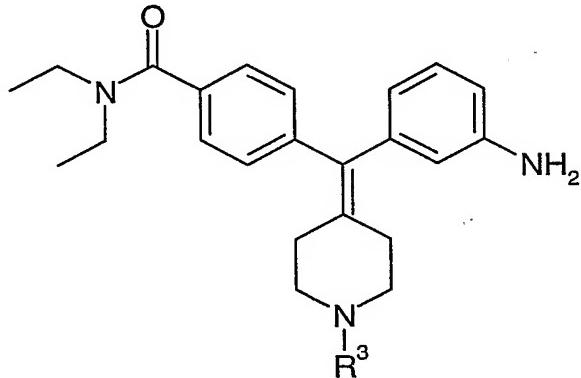
In a further aspect, the present invention provides a method of preparing a compound of formula I.

In one embodiment, the present invention provides a process for preparing a compound of formula III,

III

comprising:

reacting a compound of formula II,



15

II

with R⁹-CHO in the presence of a reducing agent to form the compound of formula III,

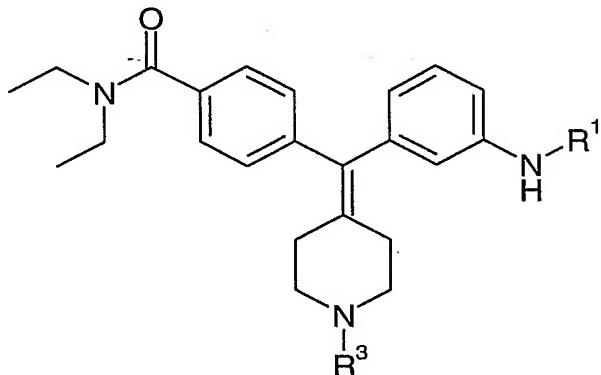
wherein

R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from

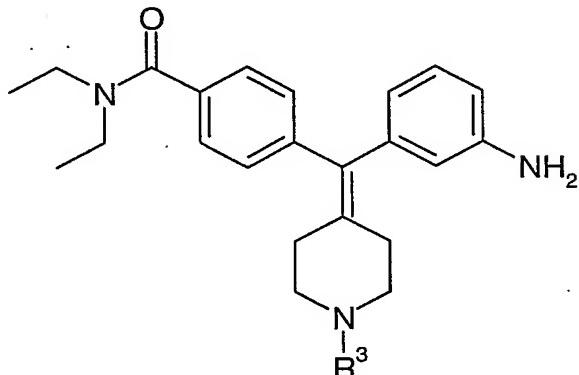
5 C_{1-4} alkyl, halogen, - CF_3 , -OH, C_{1-3} alkoxy, phenoxy and halogen; and

R^3 is selected from C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, - NO_2 , - CF_3 , C_{1-6} alkoxy and halogen.

10 In another embodiment, the present invention provides a process for preparing a compound of formula IV,

IV

comprising: reacting a compound of formula II,



15

II

with R^1-X to form the compound of formula IV,

wherein

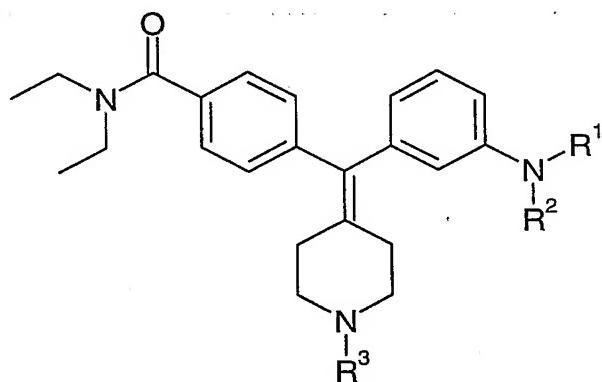
X is halogen;

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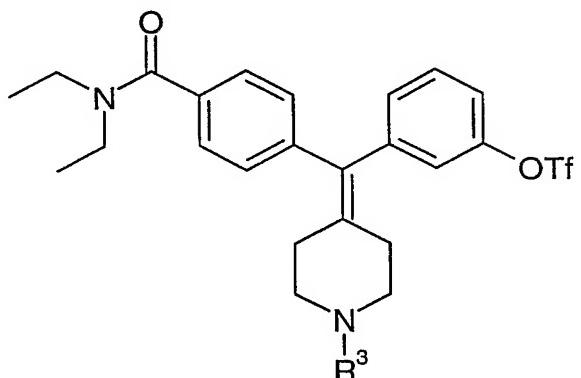
R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-4} alkyl, halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, phenoxy, and halogen; and

R^3 is selected from C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy and halogen.

In another embodiment, the present invention provides a process for preparing a compound of formula I,

I

comprising: reacting a compound of formula V,

V

with R^1R^2NH to form the compound of formula I,
wherein

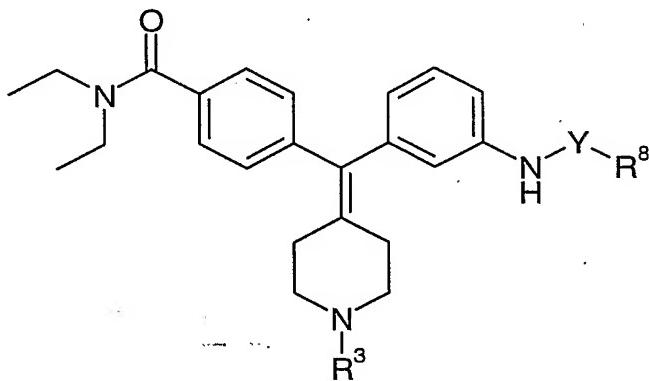
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R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-4} alkyl, halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, phenoxy, and halogen;

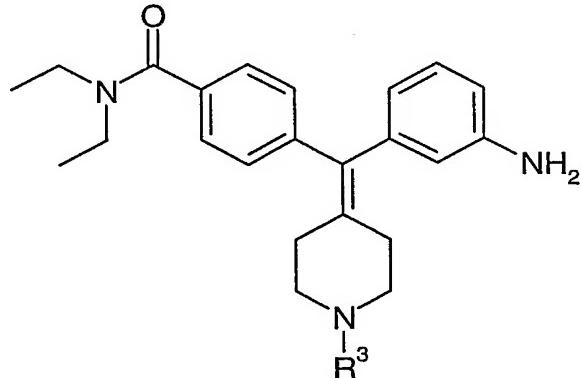
R^2 is selected from -H and C_{1-6} alkyl optionally substituted with one or more groups selected from halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, and halogen, or R^1 and R^2 are C_{1-3} alkylene that together form a portion of a ring; and

R^3 is selected from C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy and halogen.

In another embodiment, the present invention provides a process for preparing a compound of formula VI,

VI

comprising: reacting a compound of formula VII,



20

VII

with R^8 -Y-X or R^8 -Y-O-Y-R⁸ to form the compound of formula VI:

wherein

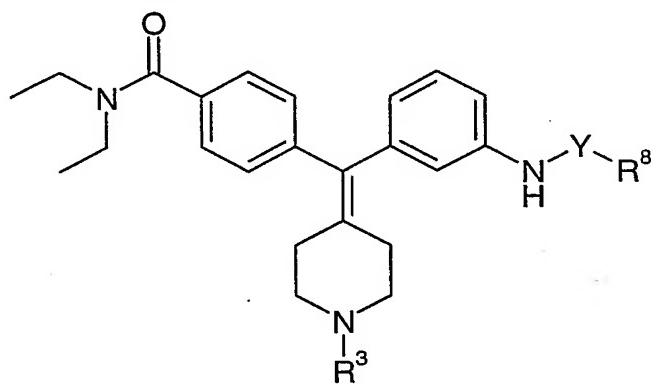
X is halogen;

5 Y is selected from -C(=O)- and -S(=O)₂;

R^8 is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

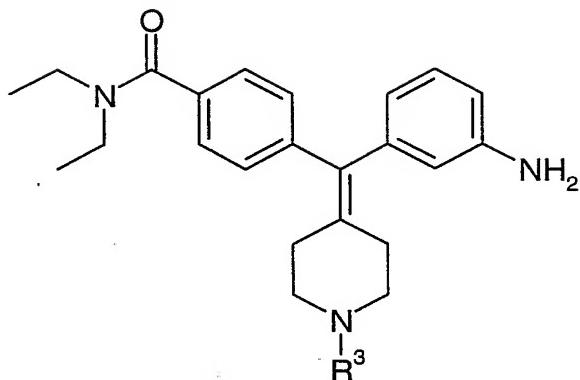
10 R^3 is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

15 In a further embodiment, the present invention provides a process for preparing a compound of formula VIII,

VIII

comprising: reacting a compound of formula VII,

21



VII

with R^8 -Z to form the compound of formula VIII:

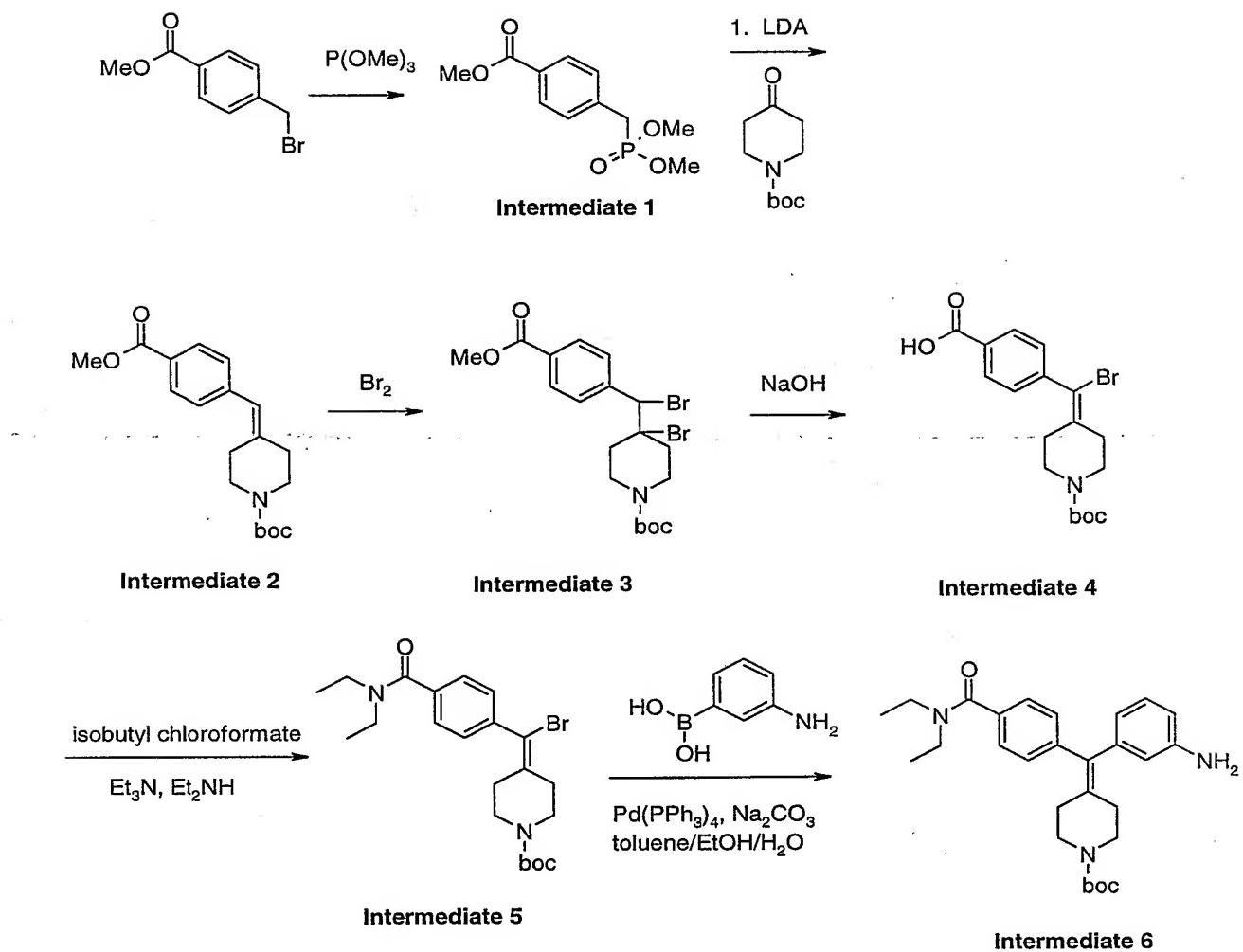
wherein

- 5 Z is selected from -NCO and -NCS;
- Y is selected from -C(=O)NH- and -C(=S)NH-;
- R^8 is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl,
- 10 C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and
- R^3 is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

Particularly, the compounds of the present invention can be prepared according to the synthetic routes as exemplified in Schemes 1-14.

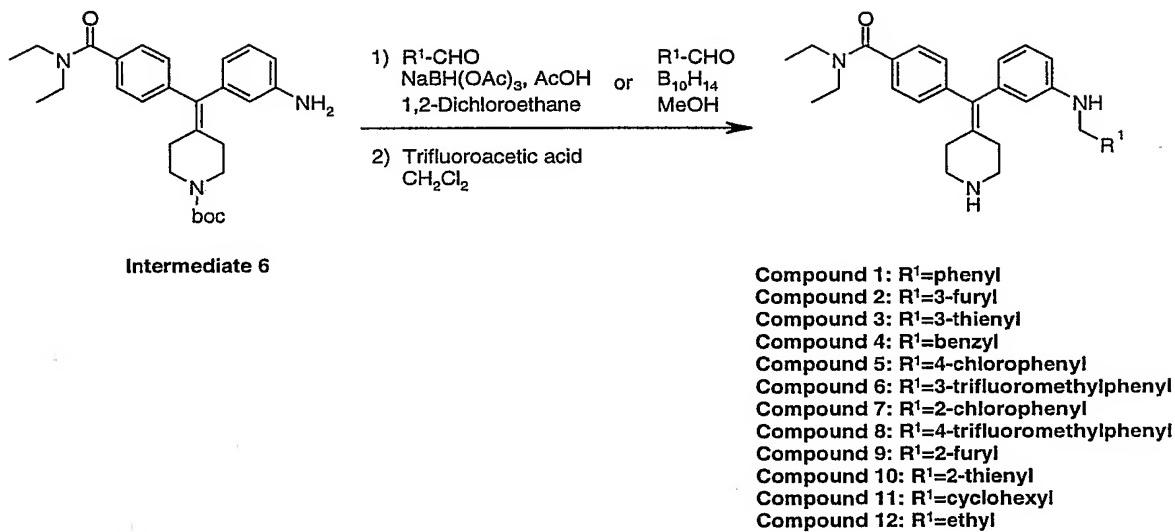
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Scheme 1

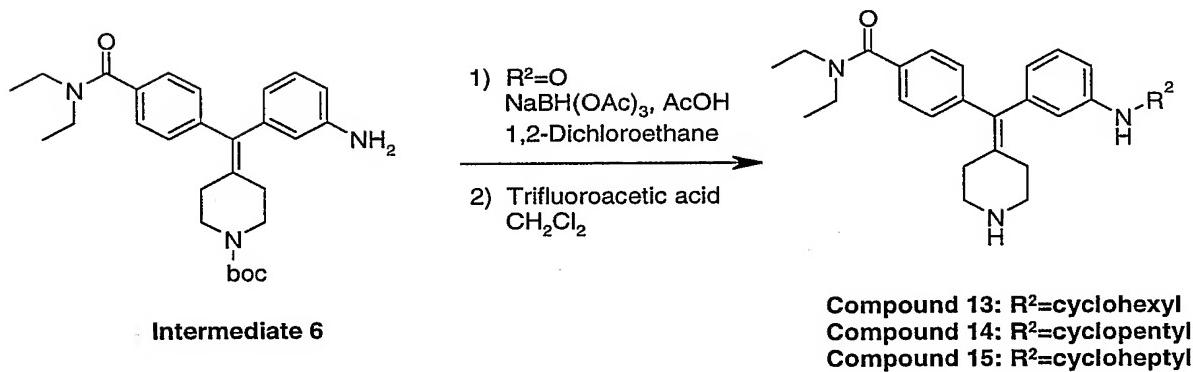


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Scheme 2

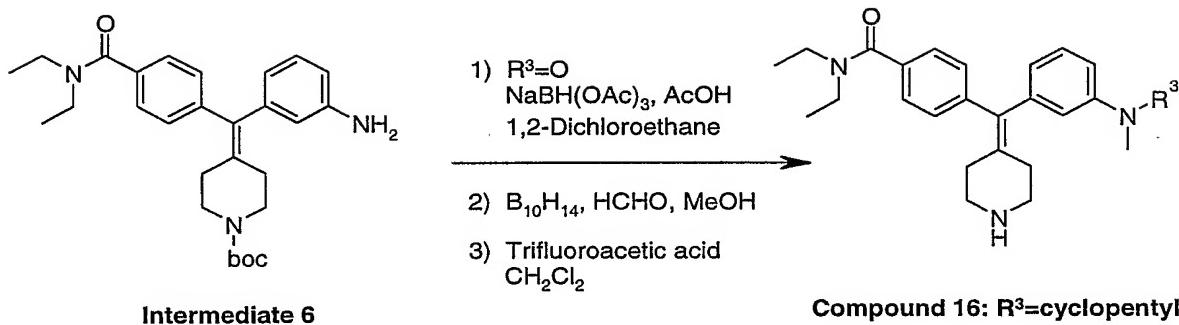


Scheme 3

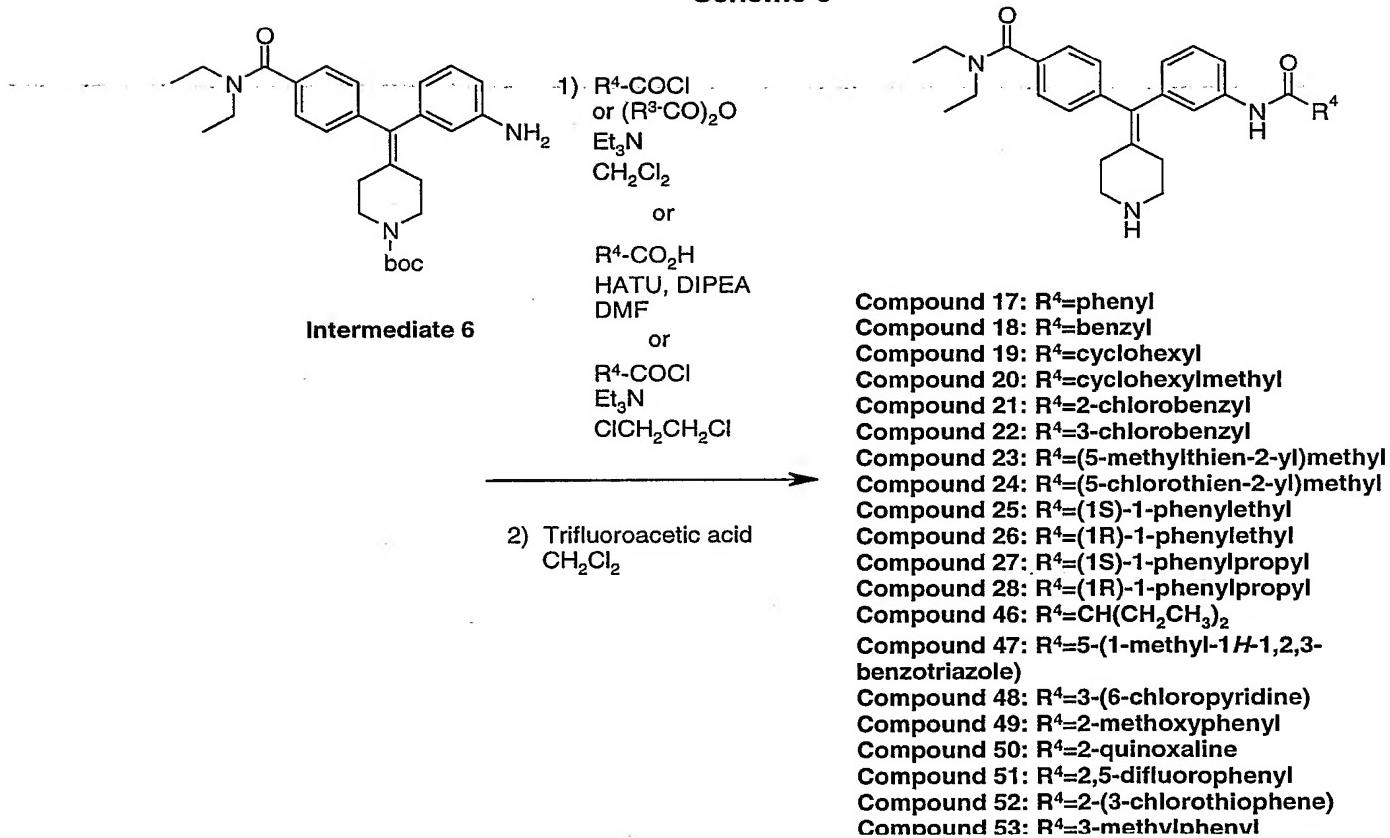


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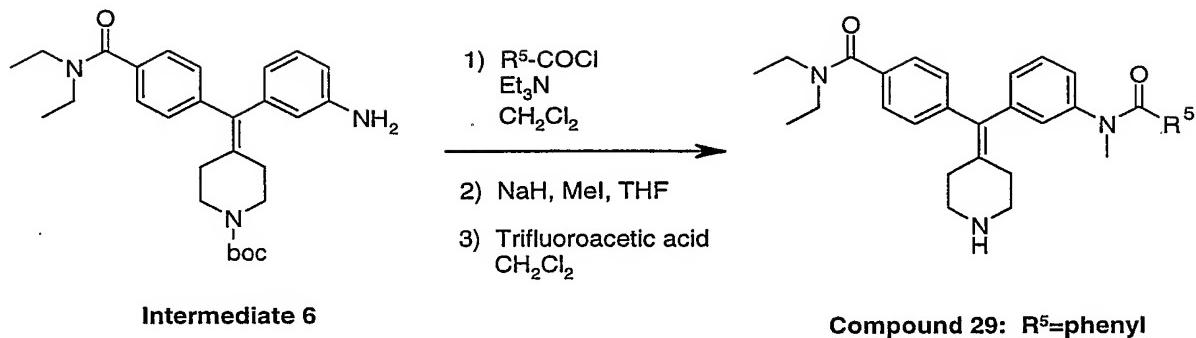
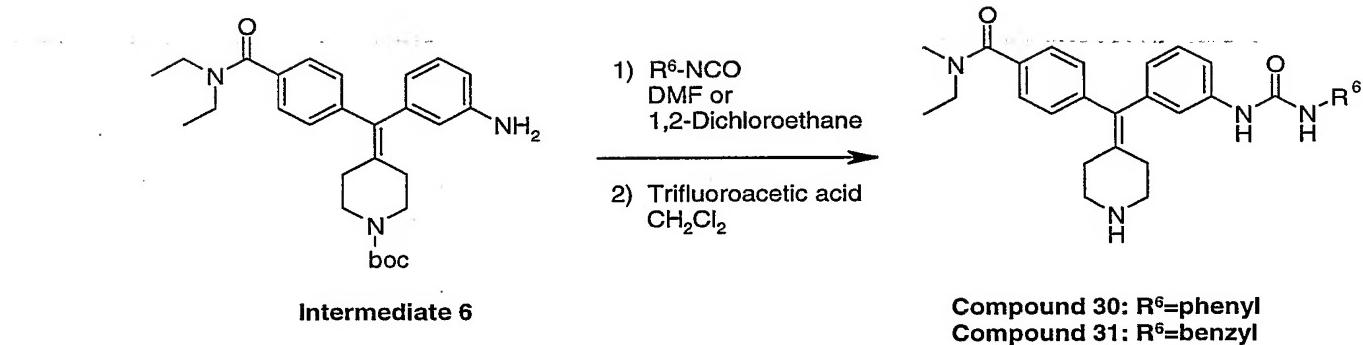
Scheme 4



Scheme 5

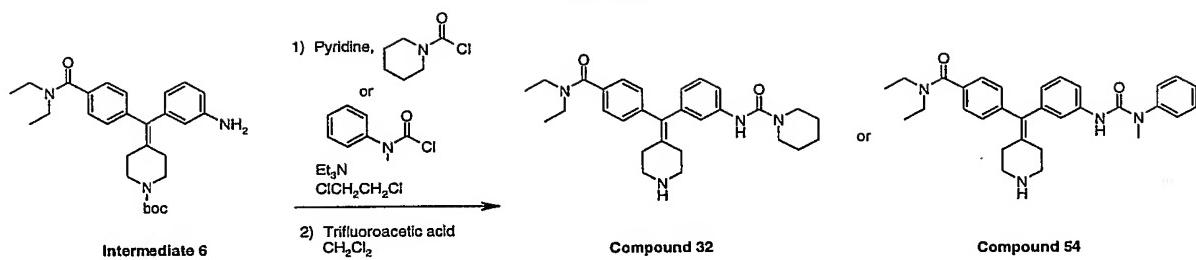
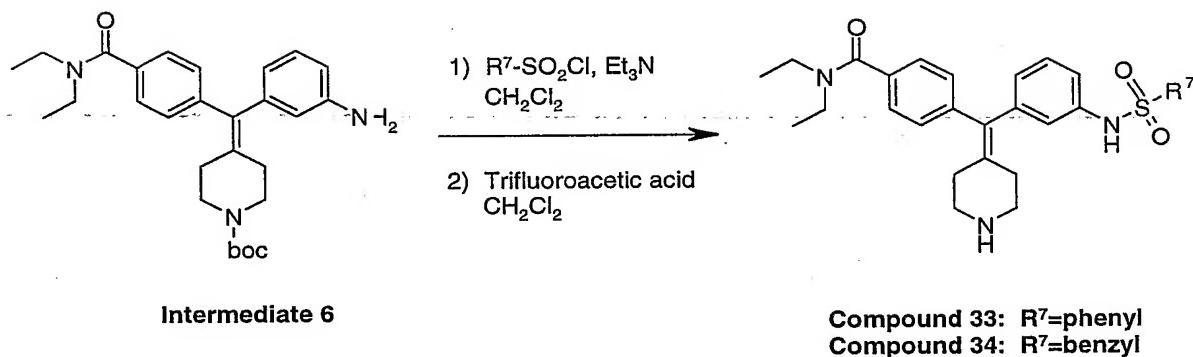


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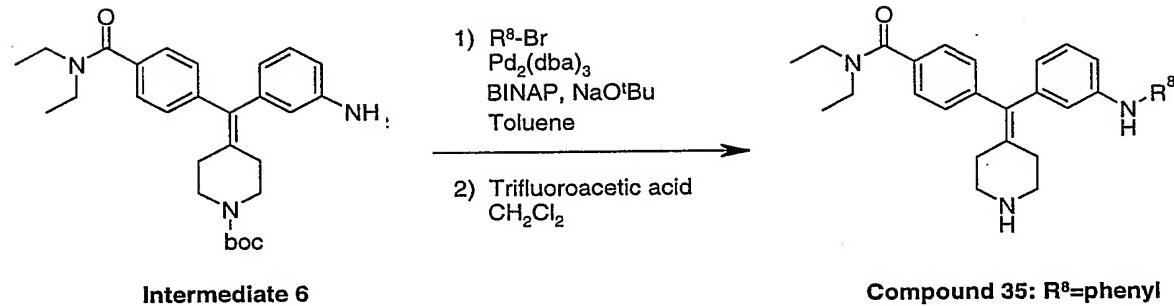
Scheme 6**Scheme 7**

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Scheme 8

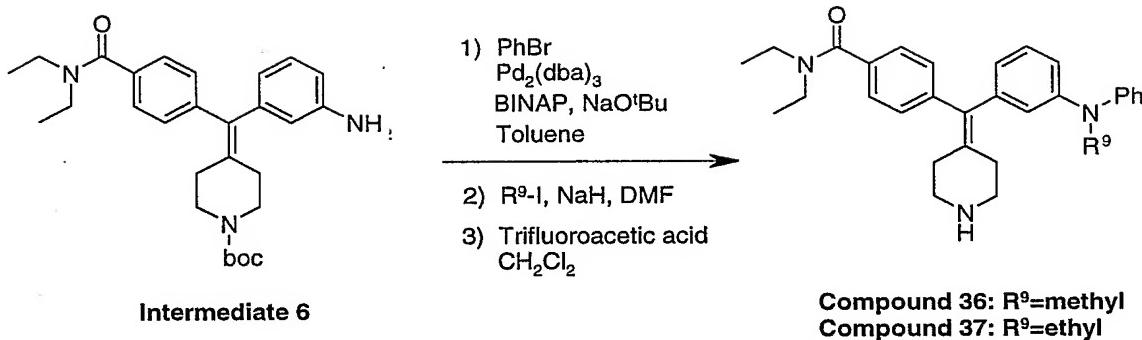
**Scheme 9**

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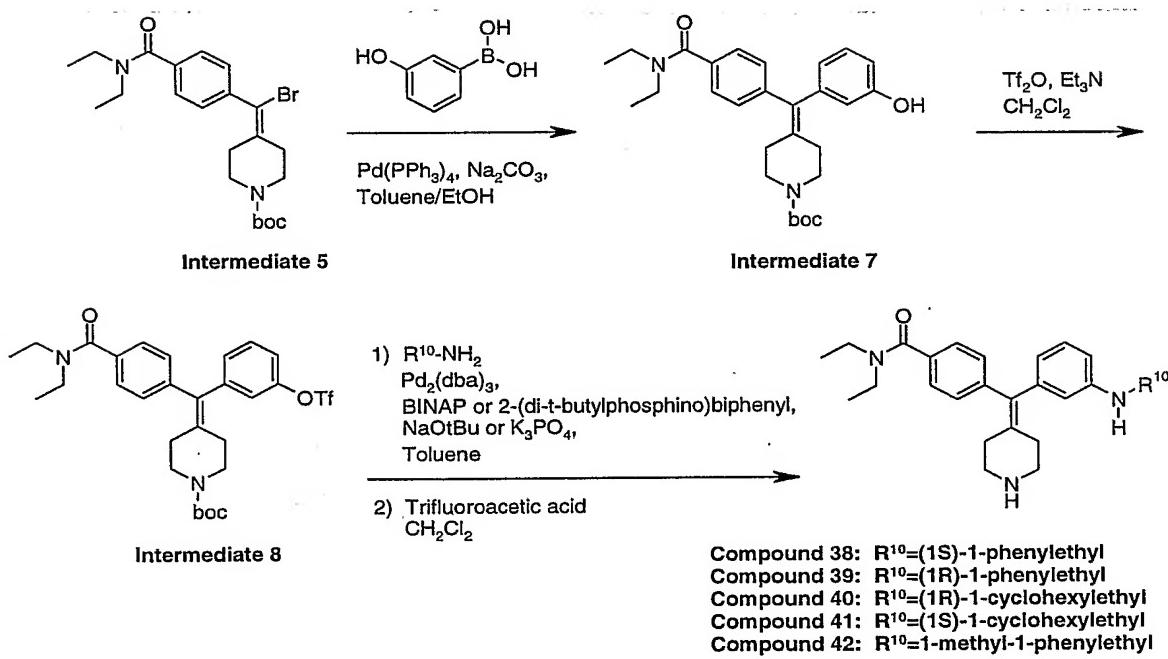
Scheme 10

27

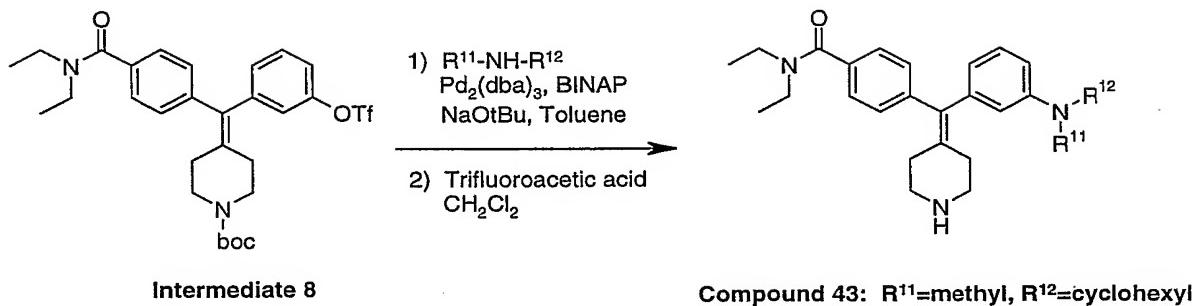
Scheme 11



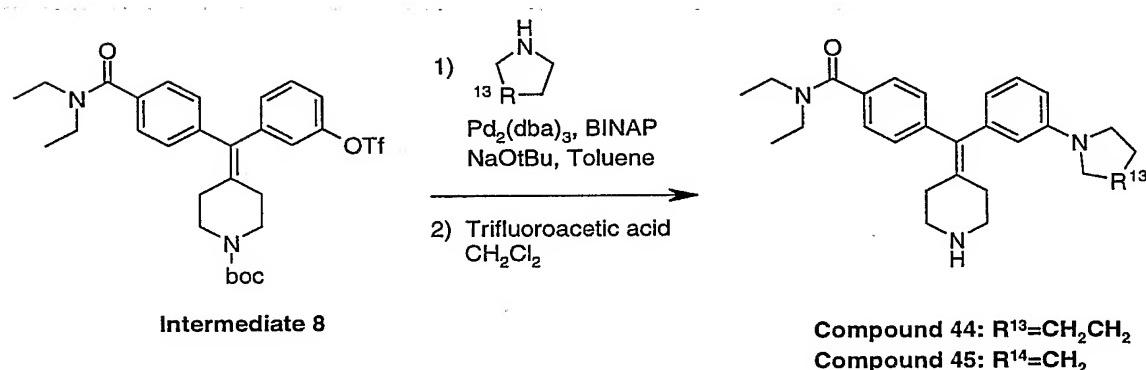
Scheme 12



Scheme 13



Scheme 14



5

BIOLOGICAL EVALUATION

The compounds of the invention are found to be active towards δ receptors in warm-blooded animal, e.g., human. Particularly the compounds of the invention are found to be effective δ receptor ligands. *In vitro* assays, *infra*, demonstrate these surprising activities, especially with regard to agonists potency and efficacy as demonstrated in the rat brain functional assay and/or the human δ receptor functional assay (low). This feature may be related to *in vivo* activity and may not be linearly correlated with binding affinity. In these *in vitro* assays, a compound is tested for their activity toward δ receptors and IC_{50} is obtained to determine the selective activity for a particular compound towards δ receptors. In the current context, IC_{50}

10
15

generally refers to the concentration of the compound at which 50% displacement of a standard radioactive δ receptor ligand has been observed.

The activities of the compound towards κ and μ receptors are also measured in a similar assay.

5 In vitro model

Cell culture

Human 293S cells expressing cloned human κ , δ and μ receptors and neomycin resistance are grown in suspension at 37°C and 5% CO₂ in shaker flasks containing calcium-free DMEM10% FBS, 5% BCS, 0.1% Pluronic F-68, and 600 µg/ml geneticin.

Rat brains are weighed and rinsed in ice-cold PBS (containing 2.5mM EDTA, pH 7.4). The brains are homogenized with a polytron for 30 sec (rat) in ice-cold lysis buffer (50mM Tris, pH 7.0, 2.5mM EDTA, with phenylmethylsulfonyl fluoride added just prior use to 0.5MmM from a 0.5M stock in DMSO:ethanol).

15 Membrane preparation

Cells are pelleted and resuspended in lysis buffer (50 mM Tris, pH 7.0, 2.5 mM EDTA, with PMSF added just prior to use to 0.1 mM from a 0.1 M stock in ethanol), incubated on ice for 15 min, then homogenized with a polytron for 30 sec. The suspension is spun at 1000g (max) for 10 min at 4°C. The supernatant is saved on ice and the pellets resuspended and spun as before. The supernatants from both spins are combined and spun at 46,000 g(max) for 30 min. The pellets are resuspended in cold Tris buffer (50 mM Tris/Cl, pH 7.0) and spun again. The final pellets are resuspended in membrane buffer (50 mM Tris, 0.32 M sucrose, pH 7.0). Aliquots (1 ml) in polypropylene tubes are frozen in dry ice/ethanol and stored at -70°C until use.

20 The protein concentrations are determined by a modified Lowry assay with sodium dodecyl sulfate.

Binding assays

Membranes are thawed at 37°C, cooled on ice, passed 3 times through a 25-gauge needle, and diluted into binding buffer (50 mM Tris, 3 mM MgCl₂, 1 mg/ml BSA (Sigma A-7888), pH 7.4, which is stored at 4°C after filtration through a 0.22 m

filter, and to which has been freshly added 5 µg/ml aprotinin, 10 µM bestatin, 10 µM diprotin A, no DTT). Aliquots of 100 µl are added to iced 12x75 mm polypropylene tubes containing 100 µl of the appropriate radioligand and 100 µl of test compound at various concentrations. Total (TB) and nonspecific (NS) binding are determined in
5 the absence and presence of 10 µM naloxone respectively. The tubes are vortexed and incubated at 25°C for 60-75 min, after which time the contents are rapidly vacuum-filtered and washed with about 12 ml/tube iced wash buffer (50 mM Tris, pH 7.0, 3 mM MgCl₂) through GF/B filters (Whatman) presoaked for at least 2h in 0.1% polyethyleneimine. The radioactivity (dpm) retained on the filters is measured with a
10 beta counter after soaking the filters for at least 12h in minivials containing 6-7 ml scintillation fluid. If the assay is set up in 96-place deep well plates, the filtration is over 96-place PEI-soaked unifilters, which are washed with 3 x 1 ml wash buffer, and dried in an oven at 55°C for 2h. The filter plates are counted in a TopCount (Packard) after adding 50 µl MS-20 scintillation fluid/well.

15 Functional Assays

The agonist activity of the compounds is measured by determining the degree to which the compounds receptor complex activates the binding of GTP to G-proteins to which the receptors are coupled. In the GTP binding assay, GTP[γ]³⁵S is combined with test compounds and membranes from HEK-293S cells expressing the cloned
20 human opioid receptors or from homogenised rat and mouse brain. Agonists stimulate GTP[γ]³⁵S binding in these membranes. The EC₅₀ and E_{max} values of compounds are determined from dose-response curves. Right shifts of the dose response curve by the delta antagonist naltrindole are performed to verify that agonist activity is mediated through delta receptors. For human δ receptor functional assays, EC₅₀ (low) is
25 measured when the human δ receptors used in the assay were expressed at lower levels in comparison with those used in determining EC₅₀ (high). The E_{max} values were determined in relation to the standard δ agonist SNC80, i.e., higher than 100% is a compound that have better efficacy than SNC80.

Procedure for rat brain GTP

30 Rat brain membranes are thawed at 37°C, passed 3 times through a 25-gauge blunt-end needle and diluted in the GTPγS binding (50 mM Hepes, 20 mM NaOH,

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100 mM NaCl, 1 mM EDTA, 5 mM MgCl₂, pH 7.4, Add fresh: 1 mM DTT, 0.1% BSA). 120 μ M GDP final is added membranes dilutions. The EC₅₀ and Emax of compounds are evaluated from 10-point dose-response curves done in 300 μ l with the appropriate amount of membrane protein (20 μ g/well) and 100000-130000 dpm of GTP γ ³⁵S per well (0.11 -0.14nM). The basal and maximal stimulated binding are determined in absence and presence of 3 μ M SNC-80

Data analysis

The specific binding (SB) was calculated as TB-NS, and the SB in the presence of various test compounds was expressed as percentage of control SB. Values of IC₅₀ and Hill coefficient (n_H) for ligands in displacing specifically bound radioligand were calculated from logit plots or curve fitting programs such as Ligand, GraphPad Prism, SigmaPlot, or ReceptorFit. Values of K_i were calculated from the Cheng-Prusoff equation. Mean \pm S.E.M. values of IC₅₀, K_i and n_H were reported for ligands tested in at least three displacement curves.

Based on the above testing protocols, we find that the compounds of the present invention and some of the intermediates used in the preparation thereof are active toward human δ receptors. Generally, the IC₅₀ towards human δ receptor for most compounds of the present invention is in the range of 0.14 nM – 31.2 nM. The EC₅₀ and %E_{max} towards human δ receptor for these compounds are generally in the range of 2.11 nM -390 nM and 89 – 118, respectively. The IC₅₀ towards human κ and μ receptors for the compounds of the invention is generally in the ranges of 36 nM- 9680 nM and 3 nM – 5975 nM, respectively.

Receptor Saturation Experiments

Radioligand K δ values are determined by performing the binding assays on cell membranes with the appropriate radioligands at concentrations ranging from 0.2 to 5 times the estimated K δ (up to 10 times if amounts of radioligand required are feasible). The specific radioligand binding is expressed as pmole/mg membrane protein. Values of K δ and B_{max} from individual experiments are obtained from

nonlinear fits of specifically bound (B) vs. nM free (F) radioligand from individual according to a one-site model.

Determination Of Mechano-Alloodynia Using Von Frey Testing

Testing is performed between 08:00 and 16:00h using the method described
5 by Chaplan et al. (1994). Rats are placed in Plexiglas cages on top of a wire mesh bottom which allows access to the paw, and are left to habituate for 10-15 min. The area tested is the mid-plantar left hind paw, avoiding the less sensitive foot pads. The paw is touched with a series of 8 Von Frey hairs with logarithmically incremental stiffness (0.41, 0.69, 1.20, 2.04, 3.63, 5.50, 8.51, and 15.14 grams; Stoelting, Ill,
10 USA). The von Frey hair is applied from underneath the mesh floor perpendicular to the plantar surface with sufficient force to cause a slight buckling against the paw, and held for approximately 6-8 seconds. A positive response is noted if the paw is sharply withdrawn. Flinching immediately upon removal of the hair is also considered a positive response. Ambulation is considered an ambiguous response, and in such
15 cases the stimulus is repeated.

Testing Protocol

The animals are tested on postoperative day 1 for the FCA-treated group. The 50% withdrawal threshold is determined using the up-down method of Dixon (1980). Testing is started with the 2.04 g hair, in the middle of the series. Stimuli are always
20 presented in a consecutive way, whether ascending or descending. In the absence of a paw withdrawal response to the initially selected hair, a stronger stimulus is presented; in the event of paw withdrawal, the next weaker stimulus is chosen. Optimal threshold calculation by this method requires 6 responses in the immediate vicinity of the 50% threshold, and counting of these 6 responses begins when the first
25 change in response occurs, e.g. the threshold is first crossed. In cases where thresholds fall outside the range of stimuli, values of 15.14 (normal sensitivity) or 0.41 (maximally allodynic) are respectively assigned. The resulting pattern of positive and negative responses is tabulated using the convention, X = no withdrawal; O = withdrawal, and the 50% withdrawal threshold is interpolated using the formula:

$$30 \quad 50\% \text{ g threshold} = 10^{(X_f + k\delta)} / 10,000$$

where X_f = value of the last von Frey hair used (log units); k = tabular value (from Chaplan et al. (1994)) for the pattern of positive / negative responses; and δ = mean difference between stimuli (log units). Here $\delta = 0.224$.

Von Frey thresholds are converted to percent of maximum possible effect (% MPE), according to Chaplan et al. 1994. The following equation is used to compute % MPE:

$$\% \text{ MPE} = \frac{\text{Drug treated threshold (g)} - \text{allodynia threshold (g)}}{\text{Control threshold (g)} - \text{allodynia threshold (g)}} \times 100$$

Administration Of Test Substance

Rats are injected (subcutaneously, intraperitoneally, intravenously or orally) with a test substance prior to von Frey testing, the time between administration of test compound and the von Frey test varies depending upon the nature of the test compound.

Writhing Test

Acetic acid will bring abdominal contractions when administered intraperitoneally in mice. These will then extend their body in a typical pattern. When analgesic drugs are administered, this described movement is less frequently observed and the drug selected as a potential good candidate.

A complete and typical Writhing reflex is considered only when the following elements are present: the animal is not in movement; the lower back is slightly depressed; the plantar aspect of *both* paws is observable. In this assay, compounds of the present invention demonstrate significant inhibition of writhing responses after oral dosing of 1-100 $\mu\text{mol/kg}$.

(i) Solutions preparation

Acetic acid (AcOH): 120 μL of Acetic Acid is added to 19.88 ml of distilled water in order to obtain a final volume of 20 ml with a final concentration of 0.6% AcOH. The solution is then mixed (vortex) and ready for injection.

Compound (drug): Each compound is prepared and dissolved in the most suitable vehicle according to standard procedures.

(ii) Solutions administration

The compound (drug) is administered orally, intraperitoneally (i.p.) ,
subcutaneously (s.c.) or intravenously (i.v.)) at 10 ml/kg (considering the average
5 mice body weight) 20, 30 or 40 minutes (according to the class of compound and its
characteristics) prior to testing. When the compound is delivered centrally:
Intraventricularly (i.c.v.) or intrathecally (i.t.) a volume of 5 µL is administered.

The AcOH is administered intraperitoneally (i.p.) in two sites at 10 ml/kg.
(considering the average mice body weight) immediately prior to testing.

10

(iii) Testing

The animal (mouse) is observed for a period of 20 minutes and the number of
occasions (Writhing reflex) noted and compiled at the end of the experiment. Mice are
kept in individual "shoe box" cages with contact bedding. A total of 4 mice are
15 usually observed at the same time: one control and three doses of drug.

For the anxiety and anxiety-like indications, efficacy has been established in
the geller-seifter conflict test in the rat.

20 For the functional gastrointestinal disorder indication, efficacy can be
established in the assay described by Coutinho SV *et al*, in American Journal of
Physiology - Gastrointestinal & Liver Physiology. 282(2):G307-16, 2002 Feb, in the
rat.

ADDITIONAL IN VIVO TESTING PROTOCOLS

Subjects and housing

Naïve male Sprague Dawley rats (175-200g) are housed in groups of 5 in a
25 temperature controlled room (22°C, 40-70% humidity, 12-h light/dark). Experiments
are performed during the light phase of the cycle. Animals have food and water ad
libitum and are sacrificed immediately after data acquisition.

Sample

Compound (Drug) testing includes groups of rats that do not receive any
30 treatment and others that are treated with E. coli lipopolysaccharide(LPS). For the
LPS-treated experiment, four groups are injected with LPS, one of the four groups is

then vehicle-treated whilst the other three groups are injected with the drug and its vehicle. A second set of experiments are conducted involving five groups of rats; all of which receive no LPS treatment. The naïve group receives no compound (drug) or vehicle; the other four groups are treated with vehicle with or without drug. These are 5 performed to determine anxiolytic or sedative effects of drugs which can contribute to a reduction in USV.

Administration of LPS

Rats are allowed to habituate in the experimental laboratory for 15-20 min prior to treatment. Inflammation is induced by administration of LPS (endotoxin of 10 gram-negative E. coli bacteria serotype 0111:B4, Sigma). LPS (2.4 μ g) is injected 15 intracerebro-ventricularly (i.c.v.), in a volume of 10 μ l, using standard stereotaxic surgical techniques under isoflurane anaesthesia. The skin between the ears is pushed rostrally and a longitudinal incision of about 1cm is made to expose the skull surface. The puncture site is determined by the coordinates: 0.8 mm posterior to the bregma, 20 1.5 mm lateral (left) to the lambda (sagittal suture), and 5 mm below the surface of the skull (vertical) in the lateral ventricle. LPS is injected via a sterile stainless steel needle (26-G 3/8) of 5 mm long attached to a 100- μ l Hamilton syringe by polyethylene tubing (PE20; 10-15 cm). A 4 mm stopper made from a cut needle (20-G) is placed over and secured to the 26-G needle by silicone glue to create the desired 25 5mm depth.

Following the injection of LPS, the needle remains in place for an additional 10 s to allow diffusion of the compound, then is removed. The incision is closed, and the rat is returned to its original cage and allowed to rest for a minimum of 3.5h prior to testing.

25 Experimental setup for air-puff stimulation

The rats remains in the experimental laboratory following LPS injection and compound (drug) administration. At the time of testing all rats are removed and placed outside the laboratory. One rat at a time is brought into the testing laboratory and placed in a clear box (9 \times 9 \times 18 cm) which is then placed in a sound-attenuating 30 ventilated cubicle measuring 62(w) \times 35(d) \times 46(h) cm (BRS/LVE, Div. Tech-Serv Inc). The delivery of air-puffs, through an air output nozzle of 0.32 cm, is controlled

by a system (AirStim, San Diego Instruments) capable of delivering puffs of air of fixed duration (0.2 s) and fixed intensity with a frequency of 1 puff per 10s. A maximum of 10 puffs are administered, or until vocalisation starts, whichever comes first. The first air puff marks the start of recording.

5 Experimental setup for and ultrasound recording

The vocalisations are recorded for 10 minutes using microphones (G.R.A.S. sound and vibrations, Vedbaek, Denmark) placed inside each cubicle and controlled by LMS (LMS CADA-X 3.5B, Data Acquisition Monitor, Troy, Michigan) software. The frequencies between 0 and 32000Hz are recorded, saved and analysed by the 10 same software (LMS CADA-X 3.5B, Time Data Processing Monitor and UPA (User Programming and Analysis)).

Compounds (Drugs)

All compounds (drugs) are pH-adjusted between 6.5 and 7.5 and administered at a volume of 4 ml/kg. Following compound (drug) administration, animals are 15 returned to their original cages until time of testing.

Analysis

The recording is run through a series of statistical and Fourier analyses to filter (between 20-24kHz) and to calculate the parameters of interest. The data are expressed as the mean \pm SEM. Statistical significance is assessed using T-test for 20 comparison between naive and LPS-treated rats, and one way ANOVA followed by Dunnett's multiple comparison test (post-hoc) for drug effectiveness. A difference between groups is considered significant with a minimum p value of ≤ 0.05 . Experiments are repeated a minimum of two times.

25 EXAMPLES

The invention will further be described in more detail by the following Examples which describe methods whereby compounds of the present invention may be prepared, purified, analyzed and biologically tested, and which are not to be construed as limiting the invention.

INTERMEDIATE 1

A mixture of 4-(bromomethyl)benzoic acid, methyl ester (11.2 g, 49 mmol) and trimethyl phosphite (25 mL) was refluxed under N₂ for 5 hrs. Excess trimethyl phosphite was removed by co-distillation with toluene to give INTERMEDIATE 1 in quantitative yield. ¹H NMR (CDCl₃) δ 3.20 (d, 2H, J=22 Hz, CH₂), 3.68 (d, 3H 10.8 Hz, OCH₃), 3.78 (d, 3H, 11.2 Hz, OCH₃), 3.91 (s, 3H, OCH₃), 7.38 (m, 2H, Ar-H), 8.00 (d, 2H, J=8 Hz, Ar-H).

INTERMEDIATE 2: 4-(4-Methoxycarbonyl-benzylidene)-piperidine-1-**carboxylic acid *tert*-butyl ester**

To a solution of INTERMEDIATE 1 in dry THF (200 mL) was added dropwise lithium diisopropylamide (32.7 mL 1.5 M in hexanes, 49 mmol) at -78 °C. The reaction mixture was then allowed to warm to room temperature prior to addition of *N*-*tert*-butoxycarbonyl-4-piperidone (9.76 g, 49 mmol in 100 mL dry THF). After 12 hrs, the reaction mixture was quenched with water (300 mL) and extracted with ethyl acetate (3 x 300 mL). The combined organic phases were dried over MgSO₄ and evaporated to give a crude product, which was purified by flash chromatography to provide INTERMEDIATE 2 as a white solid (5.64 g, 35%). IR (NaCl) 3424, 2974, 2855, 1718, 1688, 1606, 1427, 1362, 1276 cm⁻¹; ¹H NMR (CDCl₃) δ 1.44 (s, 9H), 2.31 (t, J=5.5 Hz, 2H), 2.42 (t, J=5.5 Hz, 2H), 3.37 (t, J=5.5 Hz, 2H), 3.48 (t, J=5.5 Hz, 2H), 3.87 (s, 3H, OCH₃), 6.33 (s, 1H, CH), 7.20 (d J=6.7 Hz, 2H, Ar-H), 7.94 (d, J=6.7 Hz, 2H, Ar-H); ¹³C NMR (CDCl₃) δ 28.3, 29.2, 36.19, 51.9, 123.7, 127.8, 128.7, 129.4, 140.5, 142.1, 154.6, 166.8.

25 INTERMEDIATE 3: 4-Bromo-4-[bromo-(4-methoxycarbonyl-phenyl)-methyl]-piperidine-1-carboxylic acid *tert*-butyl ester

To a mixture of INTERMEDIATE 2 (5.2 g, 16 mmol) and K₂CO₃ (1.0 g) in dry dichloromethane (200 mL) was added a solution of bromine (2.9 g, 18 mmol) in 30 mL CH₂Cl₂ at 0 °C. after 1.5 hrs at room temperature, the solution after filtration of K₂CO₃ was condensed. The residue was then dissolved in ethyl acetate (200 mL), washed with water (200 mL), 0.5 M HC1 (200 mL) and brine (200 mL), and dried over MgSO₄. Removal of solvents provided a crude product, which was

recrystallized from methanol to give INTERMEDIATE 3 as a white solid (6.07 g, 78%). IR (NaCl) 3425, 2969, 1725, 1669, 1426, 1365, 1279, 1243 cm⁻¹; ¹H NMR (CDCl₃) δ 1.28 (s, 9H), 1.75 (m, 1H), 1.90 (m, 1H), 2.1 (m, 2H), 3.08 (br, 2H), 3.90 (s, 3H, OCH₃), 4.08 (br, 3H), 7.57 (d, J=8.4 Hz, 2H, Ar-H) 7.98 (d, J=8.4 Hz, 2H, Ar-H); ¹³C NMR (CDCl₃) δ 28.3, 36.6, 38.3, 40.3, 52.1, 63.2, 72.9, 129.0, 130.3, 130.4, 141.9, 154.4, 166.3.

INTERMEDIATE 4: 4-[bromo-(4-carboxy-phenyl)-methylene]-piperidine-1-carboxylic acid tert-butyl ester

A solution of INTERMEDIATE 3 (5.4 g 11 mmol) in methanol (300 mL) and 2.0 M NaOH (100 mL) was heated at 40 °C for 3 hrs. The solid was collected by filtration, and dried overnight under vacuum. The dry salt was dissolved in 40% acetonitrile/water, and was adjusted to pH 2 using concentrated HCl. INTERMEDIATE 4 (3.8 g, 87%) was isolated as a white powder by filtration. ¹H NMR (CDCl₃) δ 1.45 (s, 9H, ^tBu), 2.22 (dd, J=5.5 Hz, 6.1 Hz, 2H), 2.64 (dd, J=5.5 Hz, 6.1 Hz, 2H), 3.34 (dd, J=5.5 Hz, 6.1 Hz, 2H), 3.54 (dd, J=5.5 Hz, 6.1 Hz, 2H), 7.35 (d, J=6.7 Hz, 2H, Ar-H), 8.08 (d, J=6.7 Hz, 2H, Ar-H); ¹³C NMR (CDCl₃) δ 28.3, 31.5, 34.2, 44.0, 115.3, 128.7, 129.4, 130.2, 137.7, 145.2, 154.6, 170.3.

INTERMEDIATE 5: 4-[bromo-(4-diethylcarbamoyl-phenyl)-methylene]-piperidine-1-carboxylic acid tert-butyl ester

To a solution of INTERMEDIATE 4 (1.0 g, 2.5 mmol) in dry dichloromethane (10 mL) at - 20 °C was added isobutylchloroformate (450 mg, 3.3 mmol). After 20 min at -20 °C diethylamine (4 mL) was added and the reaction was allowed to warm to room temperature. After 1.5 hrs the solvents were evaporated and the residue was partitioned between ethyl acetate and water. The organic phase was washed with brine and dried over MgSO₄. Removal of solvents provided a crude product, which was purified by flash chromatography to give INTERMEDIATE 5 as white needles (800 mg, 73%). IR (NaCl) 3051, 2975, 1694, 1633, 1416, 1281, 1168, 1115 cm⁻¹; ¹H NMR (CDCl₃) δ 1.13 (br, 3H, CH₃), 1.22 (br, 3H, CH₃), 1.44 (s, 9H, ^tBu), 2.22 (t, J=5.5 Hz, 2H), 2.62 (t, J=5.5 Hz, 2H), 3.33 (m, 4H), 3.55 (m, 4H), 7.31 (d, J=8.0 Hz, 2H, Ar-H), 7.36 (d, J=8.0 Hz, 2H, Ar-H); ¹³C NMR (CDCl₃) δ 12.71,

14.13, 28.3, 31.5, 34.2, 39.1, 43.2, 79.7, 115.9, 126.3, 129.3, 136.8, 137.1, 140.6,
154.6, 170.5.

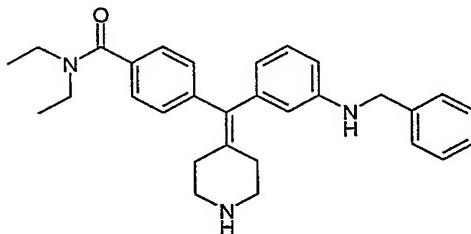
INTERMEDIATE 6: N,N-diethyl-4-(3-aminophenyl-piperidin-4-ylidene-

5 **methyl)-benzamide**

To a flask containing INTERMEDIATE 5 (5.94 g, 13.2 mmol) was added toluene (130 mL), ethanol (25 mL), 2.0 M sodium carbonate (16 mL, 32.4 mmol) and 3-aminophenylboronic acid (3.09 g, 19.9 mmol). The solution was degassed for 20 minutes, and then palladium tetrakis(triphenylphosphine) (1.53 g, 1.32 mmol) was added. The reaction mixture was heated at 90 °C overnight under N₂. The reaction was concentrated and the residue was diluted with ethyl acetate. The solution was washed with two portions of brine and the organic layer was dried (Na₂SO₄), filtered, and concentrated. The residue was purified by flash chromatography, eluting 3% methanol in dichloromethane, to yield INTERMEDIATE 6 as a colourless solid (6.12 g, 97%). (400MHz, CDCl₃) δ 1.08-1.18 (m, 3H), 1.18-1.28 (m, 3H), 2.27-2.36 (m, 4H), 3.23-3.34 (m, 2H), 3.40-3.48 (m, 2H), 3.49-3.58 (m, 2H), 3.60-3.66 (m, 2H), 6.38-6.41 (m, 1H), 6.50-6.59 (m, 2H), 7.08 (t, J = 7.60 Hz, 1H), 7.14 (d, J = 8.32 Hz, 2H), 7.30 (d, J = 8.17 Hz, 2H).

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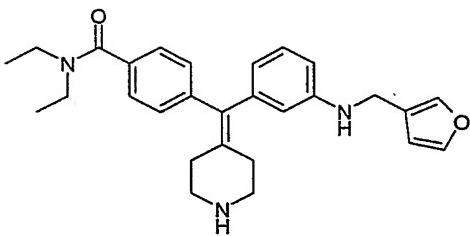
COMPOUND 1: 4-[[3-(benzylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



25 To a solution of INTERMEDIATE 6 (150 mg, 0.324 mmol) in 1,2-dichloroethane (10 mL) was added benzaldehyde (36 μL, 0.356 mmol) and sodium triacetoxyborohydride (75 mg, 0.356 mmol). The reaction was stirred at room temperature under nitrogen. After 18 hours the reaction was diluted with

dichloromethane and washed with saturated aqueous sodium bicarbonate. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried over anhydrous sodium sulfate, filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1 mL) was added. After 18 hours the solution was diluted with dichloromethane and washed with saturated aqueous sodium bicarbonate. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by reverse phase chromatography, eluting 5% to 50% acetonitrile in water containing 10 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 1 (116 mg, 53%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CDCl₃) δ 1.12-1.18 (m, 3H), 1.22-1.29 (m, 3H), 2.32 (br s, 2H), 2.46 (br s, 2H), 2.92-3.07 (m, 4H), 3.24-3.35 (m, 2H), 3.50-3.61 (m, 2H), 4.34 (s, 2H), 6.80-6.87 (m, 1H), 6.94 (s, 1H), 6.99 (d, J = 7.75 Hz, 2H), 7.16-15 7.21 (m, 1H), 7.22-7.34 (m, 9H). Found: C, 54.84; H, 5.13; N, 5.55. C₃₀H₃₅N₃O x 2.7CF₃CO₂H x 0.8H₂O has C, 54.80; H, 5.11; N, 5.42%.

COMPOUND 2: N,N-diethyl-4-[{3-[(3-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide

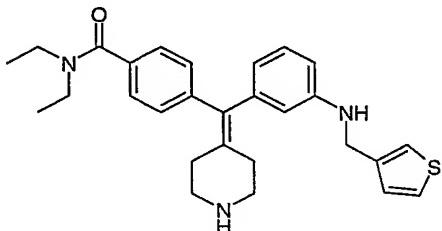


20 Using the same method as for COMPOUND 1 and using INTERMEDIATE 6 (0.315 g, 0.680 mmol) and 3-furaldehyde (67 μL, 0.77 mmol) provided COMPOUND 2. The crude material was purified by reverse phase HPLC (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid) to give COMPOUND 2 (100 mg, 22%) as its TFA salt. This material was lyophilized to produce a pale yellow solid. Purity (HPLC): >99%; ¹H NMR (400MHz, CD₃OD) δ 0.97 (br t, J = 6.25 Hz, 3H), 1.09 (br t, J = 6.25 Hz, 3H), 2.31-2.47 (m, 4H), 2.98-3.24 (m, 6H), 3.29-3.48 (m, 2H), 4.21 (s, 2H), 6.16-6.31 (m, 1H), 6.72-6.81 (m, 1H), 6.85-6.91 (m, 1H), 6.95-7.01 (m, 1H),

7.08 (d, $J = 6.44$ Hz, 2H), 7.16-7.35 (m, 5H). Found: C, 54.84; H, 5.08; N, 6.86.

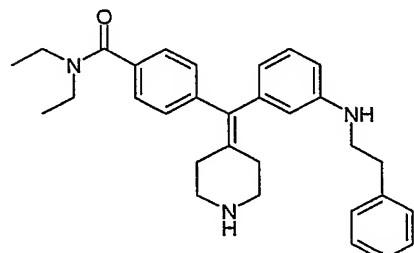
$C_{28}H_{33}N_3O_2 \times 2.1 CF_3CO_2H \times 1.1 H_2O$ has C, 55.03; H, 5.35; N, 5.98%.

COMPOUND 3: *N,N*-diethyl-4-(piperidin-4-ylidene{3-[(thien-3-ylmethyl)amino]phenyl}methyl)benzamide



Using the same method as for COMPOUND 1 and using INTERMEDIATE 6 (0.299 g, 0.646 mmol) and 3-thiophenecarboxaldehyde (62 μ L, 0.71 mmol) provided COMPOUND 3. The crude material was purified by reverse phase HPLC (gradient 10 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid) to give COMPOUND 3 (266 mg, 60%) as its TFA salt. This material was lyophilized to produce a pale yellow solid. Purity (HPLC): >96% (215 nm), >96% (254 nm), >97% (280 nm); 1H NMR (400MHz, CD₃OD) δ 0.86-1.21 (m, 6H), 2.22-2.55 (m, 4H), 3.01-3.11 (m, 4 H), 3.12-3.21 (m, 2H), 3.34-3.46 (m, 2H), 4.36 (s, 2H), 6.65-6.70 (m, 1H), 6.75-6.82 (m, 1H), 6.87-6.96 (m, 2H), 7.04-7.12 (m, 2H), 7.13-7.30 (m, 5 H). Found: C, 52.30; H, 5.02; N, 5.56. $C_{28}H_{33}N_3OS \times 2.4 CF_3CO_2H \times 1.1 H_2O$ has C, 52.31; H, 5.03; N, 5.58%.

COMPOUND 4: *N,N*-diethyl-4-[{3-[(2-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide

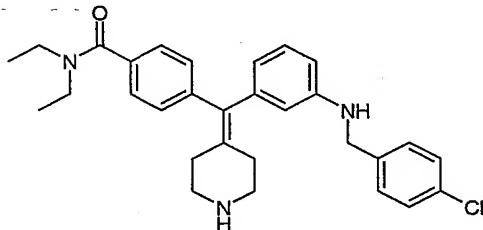


Using the same method as for COMPOUND 1 and using INTERMEDIATE 6 (0.300 g, 0.647 mmol) and phenylacetaldehyde (0.90 mL, 0.80 M solution in DCE, 0.72 mmol) provided COMPOUND 4. The crude material was purified by reverse phase

chromatography (gradient 10-45% CH₃CN in H₂O containing 0.1% trifluoroacetic acid) to give COMPOUND 4 (211 mg, 47%) as its TFA salt. This material was lyophilized to produce a colourless solid. Purity (HPLC): >97% (215 nm), >97% (254 nm), >99% (280 nm); ¹H NMR (400 MHz, CD₃OD) δ 1.09 (br t, J = 6.83 Hz, 3H), 1.22 (br t, J = 6.83 Hz, 3H), 2.55-2.65 (m, 4H), 2.84-2.92 (m, 2H), 3.22-3.29 (m, 6H), 3.38-3.44 (m, 2H), 3.48-3.57 (m, 2H), 6.74-6.77 (m, 1H), 6.81-6.86 (m, 1H), 6.89-6.94 (m, 1H), 7.17-7.24 (m, 3H), 7.25-7.33 (m, 5H), 7.36 (d, J = 8.40 Hz, 2H). Found: C, 57.43; H, 5.32; N, 5.62. C₃₁H₃₇N₃O x 2.5 CF₃CO₂H has C, 57.45; H, 5.29; N, 5.58%.

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COMPOUND 5: 4-[{3-[(4-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide

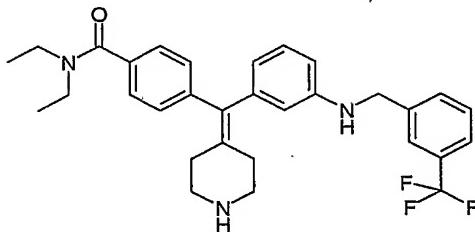


15 To a solution of INTERMEDIATE 6 (0.298 g, 0.643 mmol) in 1,2-dichloroethane (20 mL) was added 4-chlorobenzaldehyde (0.191 g, 1.36 mmol), sodium triacetoxyborohydride (0.286 g, 1.35 mmol) and glacial acetic acid (37 μL, 0.64 mmol). The reaction was stirred at room temperature under nitrogen. After 18 hours, trifluoroacetic acid (2.0 mL) was added and the reaction mixture was stirred for an additional 2 hours. Saturated aqueous sodium bicarbonate was slowly added until bubbling ceased. The organic layer was separated and the aqueous layer was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na₂SO₄), filtered, and concentrated. The residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 5 (209 mg, 45%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.03 Hz, 3H), 1.25 (br t, J = 6.64 Hz, 3H), 2.42-2.57 (m, 4 H), 3.07-3.15 (m, 2H), 3.15-3.24 (m, 2H), 3.24-

3.36 (m, 2H), 3.48-3.59 (m, 2H), 4.34 (s, 2H), 6.41-6.44 (m, 1H), 6.49-6.54 (m, 1H), 6.65-6.70 (m, 1H), 7.12 (t, J = 7.81 Hz, 1H), 7.18 (d, J = 8.01 Hz, 2H), 7.29 (s, 4H), 7.33 (d, J = 8.20 Hz, 2H) 7.22-7.34 (m, 9H). Found: C, 54.59; H, 4.97; N, 5.53.
 $C_{30}H_{34}N_3OCl \times 2.3 CF_3CO_2H \times 0.6 H_2O$ has C, 54.60; H, 4.97; N, 5.52%.

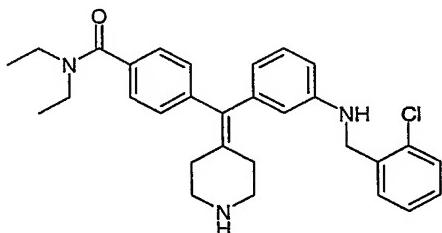
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COMPOUND 6: *N,N-diethyl-4-[piperidin-4-ylidene(3-{[3-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide*



Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.299 g, 0.645 mmol) and trifluoro-m-tolualdehyde (174 μ L, 1.30 mmol) provided COMPOUND 6 (166 mg, 34%) as a colourless solid. Purity (HPLC): >99%; 1H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.44 Hz, 3H), 1.25 (br t, J = 6.44 Hz, 3H), 2.43-2.57 (m, 4H), 3.05-3.14 (m, 2H), 3.16-3.23 (m, 2H), 3.23-3.37 (m, 2H), 3.46-3.61 (m, 2H), 4.41 (s, 2H), 6.36-6.40 (m, 1H), 6.41-6.47 (m, 1H), 6.57-6.63 (m, 1H), 7.09 (t, J = 8.01 Hz, 1H), 7.19 (d, J = 8.20 Hz, 2H), 7.31 (d, J = 8.20 Hz, 2H), 7.44-7.66 (m, 4H). Found: C, 54.84; H, 4.76; N, 5.46. $C_{31}H_{34}N_3OF_3 \times 2.2 CF_3CO_2H \times 0.2 H_2O$ has C, 54.79; H, 4.75; N, 5.41%.

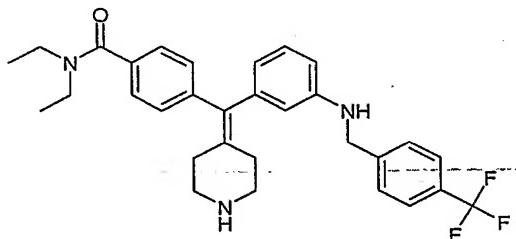
20 **COMPOUND 7:** *4-[{3-[(2-chlorobenzyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide*



Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.301 g, 0.649 mmol) and 2-chlorobenzaldehyde (153 μ L, 1.36 mmol) provided COMPOUND 7 (185 mg, 40%) as a yellow solid. Purity (HPLC): >99%; 1H NMR

(400 MHz, CD₃OD) δ 1.12 (br t, J = 6.83 Hz, 3H), 1.24 (br t, J = 6.83 Hz, 3H), 2.44-2.56 (m, 4H), 3.06-3.12 (m, 2H), 3.16-3.22 (m, 2H), 3.23-3.35 (m, 4H), 4.38-4.42 (s, 2H), 6.31-6.35 (m, 1H), 6.40-6.45 (m, 1H), 6.56-6.61 (m, 1H), 7.04-7.12 (m, 1H), 7.17-7.26 (m, 4H), 7.32 (d, J = 8.40 Hz, 2H), 7.34-7.41 (m, 2H). Found: C, 59.06; H, 5.41; N, 6.34. C₃₀H₃₄N₃OCl x 1.60 CF₃CO₂H x 0.30 H₂O has C, 59.00; H, 5.40; N, 6.22%.

COMPOUND 8: *N,N*-diethyl-4-[piperidin-4-ylidene(3-{{[4-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide

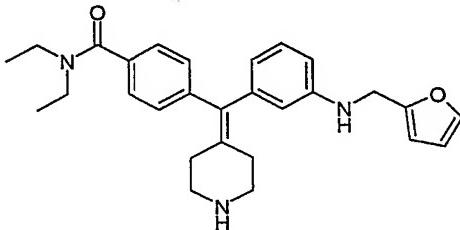


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Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.300 g, 0.649 mmol) and trifluoro-p-tolualdehyde (187 μL, 1.36 mmol) provided COMPOUND 8 (258 mg, 53%) as a colourless solid. Purity (HPLC): >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.23 Hz, 3H), 1.23 (br t, J = 7.03 Hz, 3H), 2.43-2.56 (m, 4H), 3.05-3.13 (m, 2H), 3.24-3.34 (m, 4H), 3.48-3.59 (m, 2H), 4.41 (s, 2H), 6.34-6.38 (m, 1H), 6.639-6.44 (m, 1H), 6.57 (ddd, J = 8.20, 2.34, 0.78 Hz, 1H), 7.07 (t, J = 8.01 Hz, 1H), 7.18 (d, J = 8.40 Hz, 2H), 7.31 (d, J = 8.40 Hz, 2H), 7.51 (d, J = 8.01 Hz, 2H), 7.60 (d, J = 8.01 Hz, 2H). Found: C, 55.12; H, 4.79; N, 5.46. C₃₁H₃₄N₃OF₃ x 2.1 CF₃CO₂H x 0.3 H₂O has C, 55.16; H, 4.83; N, 5.48%.

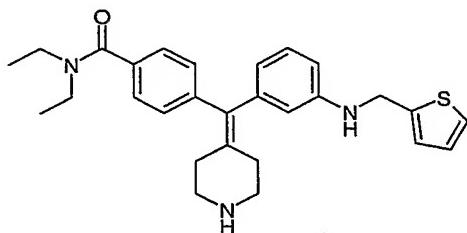
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COMPOUND 9: *N,N*-diethyl-4-[{3-[(2-furylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide



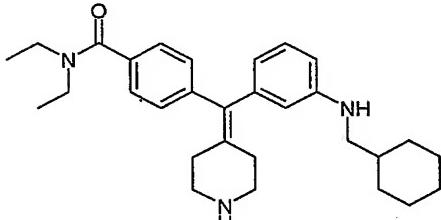
Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.297 g, 0.641 mmol) and furfural (110 μ L, 1.33 mmol) provided COMPOUND 9. The crude material was purified by reverse phase chromatography (gradient 20-80% acetonitrile in water containing 0.1% trifluoroacetic acid) to give COMPOUND 9 (100 mg, 23%) as its TFA salt. This material was lyophilized to produce a yellow solid. Purity (HPLC): >99%; 1 H NMR (400 MHz, CD₃OD) δ 1.13 (br t, J = 7.42 Hz, 3H), 1.23 (t, J = 7.42 Hz, 3H), 2.52-2.59 (m, 4H), 3.17-3.26 (m, 4H), 3.26-3.34 (m, 2H), 3.48-3.59 (m, 2H), 4.30 (s, 2H), 6.17-6.19 (m, 1H), 6.31 (dd, J = 3.12, 1.76 Hz, 1H), 6.50-6.56 (m, 2H), 6.69 (ddd, J = 8.20, 2.15, 0.98 Hz, 1H), 7.13 (t, J = 8.01 Hz, 1H), 7.24 (d, J = 8.40 Hz, 2H), 7.34 (d, J = 8.40 Hz, 2H), 7.40 (dd, J = 1.95, 0.78 Hz, 1H). Found: C, 54.91; H, 5.03; N, 5.96. C₂₈H₃₃N₃O₂ x 2.30 CF₃CO₂H x 0.40 H₂O has C, 54.91; H, 5.10; N, 5.89%.

COMPOUND 10: N,N-diethyl-4-(piperidin-4-ylidene{3-[{(thien-2-yl)methyl]amino}phenyl}methyl)benzamide



Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.289 g, 624 mmol) and 2-thiophenecarboxaldehyde (0.12 mL, 1.28 mmol) provided COMPOUND 10. The crude material was purified by reverse phase chromatography (gradient 10-60% acetonitrile in water containing 0.1% trifluoroacetic acid) to give COMPOUND 10 (239 mg, 56%) as its TFA salt. This material was lyophilized to produce a colourless solid. Purity (HPLC): >94% (215 nm), >97% (254 nm), >99% (280 nm); 1 H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.03 Hz, 3H), 1.22 (br t, J = 7.03 Hz, 3H), 2.47-2.59 (m, 4H), 3.11-3.17 (m, 2H), 3.17-3.24 (m, 2H), 3.24-3.36 (m, 2H), 3.48-3.58 (m, 2H), 4.50-4.55 (s, 2H), 6.49-6.55 (m, 2H), 6.68-6.73 (m, 1H), 6.91-6.99 (m, 2H), 7.13 (d, J = 7.23 Hz, 1H), 7.22 (d, J = 7.42 Hz, 2H), 7.24-7.27 (m, 1H), 7.33 (d, J = 7.62 Hz, 2H). Found: C, 55.30; H, 5.23; N, 6.06. C₂₈H₃₃N₃OS x 2.0 CF₃CO₂H x 0.4 H₂O has C, 55.31; H, 5.19; N, 6.05%.

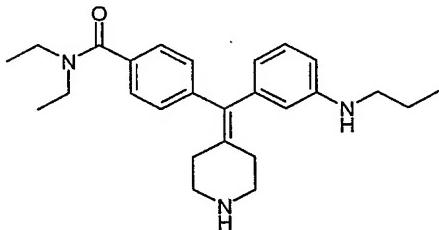
COMPOUND 11: 4-[{3-[(cyclohexylmethyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



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Using the same method as for COMPOUND 5 and using INTERMEDIATE 6 (0.198 g, 0.428 mmol) and cyclohexanecarboxaldehyde (0.10 mL, 0.83 mmol) provided COMPOUND 11 (147 mg, 50%) as a yellow solid. Purity (HPLC): >99%; ^1H NMR (400 MHz, CD₃OD) δ 0.96-1.08 (m, 2H), 1.12 (t, J = 6.83 Hz, 3H), 1.17-1.32 (m, 6H), 1.57-1.85 (m, 6H), 2.56-2.64 (m, 4H), 3.07 (d, J = 6.83 Hz, 2H), 3.22-3.35 (m, 6H), 3.49-3.58 (m, 2H), 6.90-6.93 (m, 1H), 6.95-6.99 (m, 1H), 7.02-7.07 (m, 1H), 7.27 (d, J = 8.20 Hz, 2H), 7.33-7.40 (m, 3H). Found: C, 56.22; H, 6.30; N, 5.36. C₃₀H₄₁N₃O x 2.3 CF₃CO₂H x 1.0 H₂O has C, 56.16; H, 6.17; N, 5.68%.

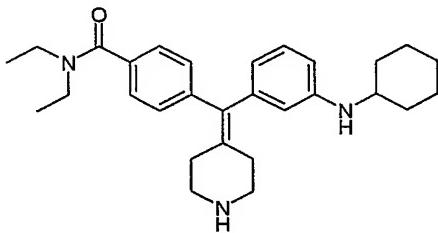
15 **COMPOUND 12: N,N-diethyl-4-{piperidin-4-ylidene[3-(propylamino)phenyl]methyl}benzamide**



To a solution of INTERMEDIATE 6 (0.299 g, 0.645 mmol) in methanol (5 mL) was added propionaldehyde (47 μ L, 0.64 mmol). The reaction was stirred for 30 minutes 20 and then decaborane (24 mg, 0.19 mmol) was added. After 2 hours, the reaction was complete. The mixture was diluted with dichloromethane and washed with saturated aqueous sodium bicarbonate. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried (Na₂SO₄), filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and

trifluoroacetic acid (1.5 mL) was added. The reaction was stirred overnight. The reaction mixture was concentrated *in vacuo* and purified by reverse phase chromatography (gradient 10-50% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 12 (107 mg, 26%) as a colourless solid. Purity (HPLC): >99%; ¹H NMR (400 MHz, CD₃OD) δ 0.99 (t, J = 7.42 Hz, 3H), 1.12 (br t, J = 6.64 Hz, 3H), 1.24 (br t, J = 6.64 Hz, 3H), 1.62-1.73 (sextet, 2H), 2.57-2.63 (m, 4H), 3.18-3.24 (m, 2H), 3.18-3.24 (m, 2H), 3.24-3.33 (m, 6H), 3.49-3.58 (m, 2H), 6.97-6.99 (m, 1H), 7.06-7.10 (m, 1H), 7.10-7.14 (m, 1H), 7.27 (d, J = 8.40 Hz, 2H), 7.37 (d, J = 8.40 Hz, 2H), 7.42 (t, J = 7.81 Hz, 1H). Found: C, 51.78; H, 4.74; N, 5.85. C₂₆H₃₅N₃O x 2.9 CF₃CO₂H has C, 51.88; H, 5.19; N, 5.71%.

COMPOUND 13: 4-[[3-(cyclohexylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



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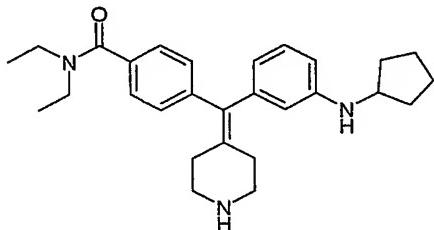
To a solution of INTERMEDIATE 6 (0.200 g, 0.432 mmol) in 1,2-dichloroethane (15 mL) was added cyclohexanone (50 μL, 0.482 mmol), sodium triacetoxyborohydride (0.138 g, 0.651 mmol) and glacial acetic acid (25 μL, 0.44 mmol). The reaction was stirred at room temperature under nitrogen. After 24 hours, trifluoroacetic acid (2.0 mL) was added and the reaction mixture was stirred for an additional 2 hours.

Saturated aqueous sodium bicarbonate was slowly added until bubbling ceased. The organic layer was separated and the aqueous layer was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na₂SO₄), filtered, and concentrated. The residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 13 (187 mg, 64%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.07-1.16 (m, 3H), 1.17-1.28 (m, 4H), 1.28-1.39 (m, 4H), 1.64-1.72 (m,

1H), 1.77-1.87 (m, 2H), 1.88-1.98 (m, 2H), 2.58-2.64 (m, 4H), 3.24-3.41 (m, 7H), 3.49-3.58 (m, 2H), 7.07-7.07 (m, 1H), 7.19-7.25 (m, 2H), 7.27 (d, $J = 8.40$ Hz, 2H), 7.38 (d, $J = 8.40$ Hz, 2H), 7.50 (t, $J = 7.81$ Hz, 1H). Found: C, 55.13; H, 5.78; N, 5.58. $C_{29}H_{39}N_3O \times 2.5 CF_3CO_2H \times 0.6 H_2O$ has C, 55.07; H, 5.80; N, 5.67%.

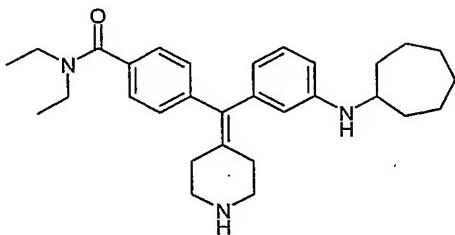
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COMPOUND 14: 4-[[3-(cyclopentylamino)phenyl](piperidin-4-ylidene)methyl]-*N,N*-diethylbenzamide

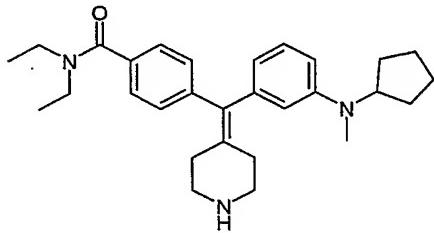


To a solution of INTERMEDIATE-6 (0.117 g, 0.252 mmol) in 1,2-dichloroethane (10 mL) was added cyclopentanone (70 μ L, 0.78 mmol), sodium triacetoxyborohydride (0.187 g, 0.882 mmol) and glacial acetic acid (30 μ L, 0.52 mmol). The reaction was stirred at room temperature under an atmosphere of nitrogen. After 24 hours, trifluoroacetic acid (1.0 mL) was added and the reaction mixture was stirred for an additional 2 hours. Saturated aqueous sodium bicarbonate was slowly added until bubbling ceased. The organic layer was separated and the aqueous layer was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na_2SO_4), filtered, and concentrated. The residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 14 (120 mg, 72%) as a colourless solid. Purity (HPLC) >99%; 1H NMR (400 MHz, CD_3OD) δ 1.09 (t, $J = 6.64$ Hz, 3H), 1.21 (t, $J = 6.64$ Hz, 3H), 1.52-1.68 (m, 4H), 1.69-1.82 (m, 2H), 1.88-2.00 (m, 2H), 2.54-2.61 (m, 4H), 3.20-3.31 (m, 6H), 3.46-3.56 (m, 2H), 3.80-3.90 (m, 1H), 6.98-7.01 (m, 1H), 7.09-7.17 (m, 2H), 7.24 (d, $J = 8.40$ Hz, 2H), 7.34 (d, $J = 8.40$ Hz, 2H), 7.43 (t, $J = 8.01$ Hz, 1H). Found: C, 55.10; H, 5.87; N, 5.65. $C_{28}H_{37}N_3O \times 2.3 CF_3CO_2H \times 1.0 H_2O$ has C, 55.00; H, 5.85; N, 5.90%.

COMPOUND 15: 4-[[3-(cycloheptylamino)phenyl](piperidin-4-ylidene)methyl]-

N,N-diethylbenzamide

Using the same method as for COMPOUND 14 and using INTERMEDIATE 6 (0.409 g, 0.883 mmol) and cycloheptanone (0.32 mL, 2.7 mmol) provided COMPOUND 15 (292 mg, 48%) as a colourless solid. Purity (HPLC): >99%; ^1H NMR (400 MHz, CD₃OD) δ 1.13 (br t, J = 7.62 Hz, 3H), 1.23 (br t, J = 7.03 Hz, 3H), 1.39-1.68 (m, 8H), 1.68-1.79 (m, 2H), 1.92-2.02 (m, 2H), 2.58-2.64 (m, 4H), 3.24-3.34 (m, 6H), 3.49-3.63 (m, 3H), 7.11-7.14 (m, 1H), 7.24-7.31 (m, 4H), 7.38 (d, J = 8.40 Hz, 2H), 7.49-7.55 (m, 1H). Found: C, 55.82; H, 5.86; N, 5.73. C₃₀H₄₁N₃O x 2.5 CF₃CO₂H x 0.4 H₂O has C, 55.91; H, 5.94; N, 5.59%.

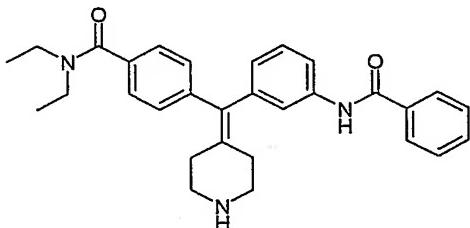
COMPOUND 16: 4-[{3-[cyclopentyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide

To a solution of INTERMEDIATE 6 (0.390 g, 0.841 mmol) in 1,2-dichloroethane (20 mL) was added cyclopentanone (0.22 mL, 2.5 mmol), sodium triacetoxyborohydride (0.570 g, 2.69 mmol) and glacial acetic acid (50 μL , 0.84 mmol). The reaction was stirred overnight at room temperature under an atmosphere of nitrogen. The mixture was washed with two portions of saturated aqueous sodium bicarbonate and one portion of brine. The organic phase was dried (Na_2SO_4), filtered, and concentrated. The residue was purified by flash chromatography, eluting 0% to 50% ethyl acetate in hexanes to yield 0.356 g of product as a colourless foam. The material was dissolved in methanol (5 mL) and formaldehyde (37% in water) (150 μL , 2.01 mmol) was added. The reaction was stirred for 30 minutes and then

decaborane (49 mg, 0.40 mmol) was added. After 18 h, the reaction mixture was concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1.5 mL) was added. The reaction was stirred for 4 hours at room temperature and then saturated aqueous sodium bicarbonate was added. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried (Na_2SO_4), filtered and concentrated. The residue was purified by reverse phase chromatography (gradient 10-50% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 16 (260 mg, 58%) as a colourless solid. Purity (HPLC) >93% (215 nm), >99% (254 nm), >99% (280 nm); ^1H NMR (400 MHz, CD_3OD) δ 1.12 (br t, $J = 6.64$ Hz, 3H), 1.24 (br t, $J = 7.62$ Hz, 3H), 1.58-1.72 (m, 4H), 1.73-1.84 (m, 2H), 1.85-2.03 (m, 2H), 2.57-2.66 (m, 4H), 3.22-3.35 (m, 9H), 3.49-3.59 (m, 2H), 4.09-4.20 (m, 1H), 7.28 (d, $J = 8.01$ Hz, 2H), 7.35-7.43 (m, 4H), 7.51-7.56 (m, 1H), 7.58-7.65 (m, 1H). Found: C, 52.68; H, 5.63; N, 5.33.

$\text{C}_{29}\text{H}_{39}\text{N}_3\text{O} \times 2.8 \text{ CF}_3\text{CO}_2\text{H} \times 1.3 \text{ H}_2\text{O}$ has C, 52.72; H, 5.68; N, 5.33%.

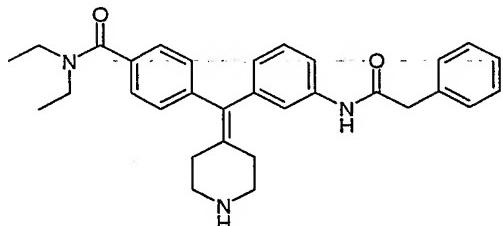
COMPOUND 17: 4-[[3-(benzoylamino)phenyl](piperidin-4-ylidene)methyl]-*N,N*-diethylbenzamide



To a solution of INTERMEDIATE 6 (0.158 g, 0.341 mmol) in dichloromethane (10 mL) was added benzoic anhydride (86 mg, 0.38 mmol) and triethylamine (0.15 mL, 1.08 mmol). The reaction was stirred for 24 h at room temperature under an atmosphere of nitrogen. The mixture was washed with saturated aqueous sodium bicarbonate. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried (Na_2SO_4), filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1.0 mL) was added. After 2 hours, the solution was concentrated and the residue was purified by reverse phase chromatography, eluting 10% to 45%

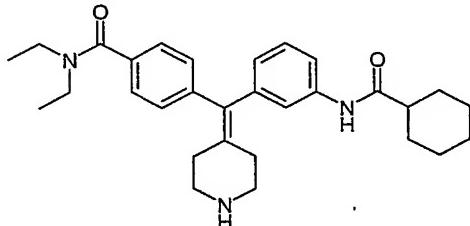
acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 17 (140 mg, 71%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.13 (br t, J = 6.83 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 2.58-2.63 (m, 2H), 2.64-2.69 (m, 2H), 3.23-3.35 (m, 6H), 3.49-3.58 (m, 2H), 6.92-6.96 (m, 1H), 7.27-7.33 (m, 2H), 7.33-7.39 (m, 3H), 7.47-7.54 (m, 3H), 7.55-7.61 (m, 1H), 7.69-7.73 (m, 1H), 7.88-7.94 (m, 2H). Found: C, 61.96; H, 5.57; N, 6.59. C₃₀H₃₃N₃O₂ x 1.4 CF₃CO₂H x 0.5 H₂O has C, 61.92; H, 5.61; N, 6.60%.

10 **COMPOUND 18: N,N-diethyl-4-[{3-[(phenylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide**



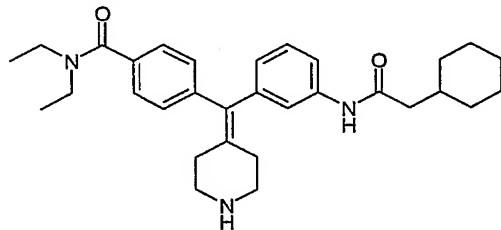
To a solution of INTERMEDIATE 6 (0.154 g, 0.332 mmol) in dichloromethane (10 mL) was added phenylacetyl chloride (48 μL, 0.36 mmol) and triethylamine (0.15 mL, 1.08 mmol). The reaction was stirred for 20 hours at room temperature under an atmosphere of nitrogen. The mixture was washed with saturated aqueous sodium bicarbonate. The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried (Na₂SO₄), filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1.0 mL) was added. After 2 hours, the solution was concentrated and the residue was purified by reverse phase chromatography, eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 18 (181 mg, 91%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.25 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.54-2.64 (m, 4H), 3.20-3.26 (m, 4H), 3.27-3.34 (m, 2H), 3.48-3.58 (m, 2H), 3.65 (s, 2H), 6.86-6.91 (m, 1H), 7.21-7.40 (m, 11H), 7.56-7.60 (m, 1H). Found: C, 59.65; H, 5.46; N, 5.91. C₃₁H₃₅N₃O₂ x 1.8 CF₃CO₂H x 0.6 H₂O has C, 59.57; H, 5.49; N, 6.02%.

COMPOUND 19: 4-[{3-[(cyclohexylcarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



5 Using the same method as for COMPOUND 18 and using INTERMEDIATE 6 (0.105 g, 0.227 mmol) and cyclohexanecarbonyl chloride (33 μ L, 0.25 mmol) provided COMPOUND 19 (110 mg, 83%) as a colourless solid. Purity (HPLC): >99%; 1 H NMR (400 MHz, CD₃OD) δ 1.12 (m, 3H), 1.18-1.41 (m, 6H), 1.49 (m, 2H), 1.72 (m, 1H), 1.83 (m, 4H), 2.33 (m, 1H), 2.60 (m, 4H), 3.22-3.34 (m, 6H), 3.53 (m, 2H), 6.87 (dt, J = 7.42, 1.56 Hz, 1H), 7.24-7.29 (m, 3H), 7.32 (m, 1H), 7.35 (d, J = 8.59 Hz, 2H), 7.57 (m, 1H). Found: C, 59.33; H, 6.31; N, 6.21. C₃₀H₃₉N₃O₂ x 1.6 CF₃CO₂H x 0.9 H₂O has C, 59.31; H, 6.36; N, 6.25%.

15 **COMPOUND 20: 4-[{3-[(cyclohexylacetyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide**

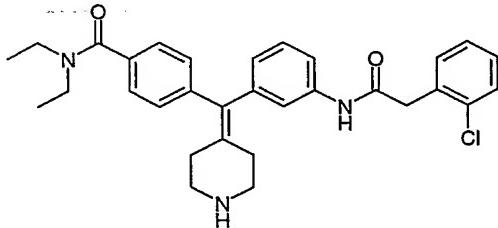


To a solution of INTERMEDIATE 6 (0.105 g, 0.226 mmol) in dry DMF (5 mL) was added cyclohexylacetic acid (50 mg, 0.35 mmol), HATU (0.122 g, 0.321 mmol), and *N,N*-diisopropylethylamine (100 μ L, 0.574 mmol). The reaction was 20 stirred overnight at room temperature under nitrogen, then concentrated *in vacuo*. The residue was dissolved in dichloromethane and washed with 2N NaOH (1x) and brine (1x). The organic phase was dried (Na₂SO₄), filtered and concentrated. The residue was dissolved in dichloromethane (5 mL) and trifluoroacetic acid (0.5 mL) was added. The reaction was stirred for 4 hours at room temperature. The solution

was concentrated and the residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as its TFA salt and was lyophilized to give COMPOUND 20 (97 mg, 71%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 0.96-1.37 (m, 11H), 1.62-1.88 (m, 6H), 2.21 (d, J = 7.03 Hz, 2H), 2.56-2.24 (m, 4H), 3.21-3.34 (m, 6H), 3.48-3.59 (m, 2H), 6.88 (dt, J = 7.42, 1.37 Hz, 1H), 7.24-7.30 (m, 3H), 7.32 (m, 1H), 7.36 (d, J = 8.40 Hz, 2H), 7.56 (m, 1H). Found: C, 60.39; H, 6.68; N, 6.10. C₃₁H₄₁N₃O₂ x 1.5 CF₃CO₂H x 1.0 H₂O has C, 60.34; H, 6.63; N, 6.21%.

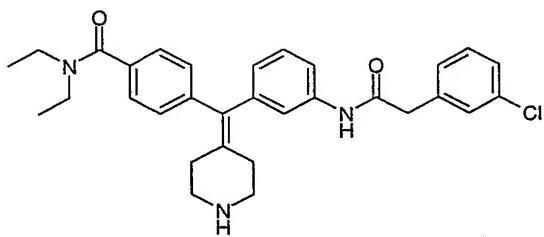
10

COMPOUND 21: 4-[(3-{[(2-chlorophenyl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.138 g, 0.298 mmol) and 2-chlorophenylacetic acid (77 mg, 0.45 mmol) provided COMPOUND 20 (115 mg, 61%) as a colourless solid. Purity (HPLC): >96% (215 nm), >96% (254 nm), >99% (280 nm); ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.83 Hz, 3H), 1.23 (br t, J = 6.83 Hz, 3H), 2.54-2.65 (m, 4H), 3.20-3.26 (m, 4H), 3.26-3.34 (m, 2H), 3.48-3.59 (m, 2H), 3.85 (s, 2H), 6.86-6.90 (m, 1H), 7.22-7.31 (m, 5H), 7.31-7.42 (m, 5H), 7.56-7.60 (m, 1H). Found: C, 59.20; H, 5.20; N, 6.08. C₃₁H₃₄N₃O₂Cl x 1.5 CF₃CO₂H x 0.1 H₂O has C, 59.28; H, 5.22; N, 6.10%.

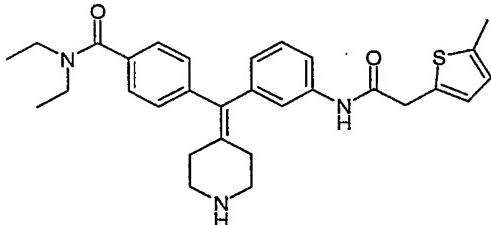
COMPOUND 22: 4-[(3-{[(3-chlorophenyl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.139 g, 0.300 mmol) and 3-chlorophenylacetic acid (77 mg, 0.45 mmol) provided COMPOUND 22 (125 mg, 66%) as a colourless solid. Purity (HPLC): >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.10 (br t, J = 6.83 Hz, 3H), 1.22 (br t, J = 6.83 Hz, 3H), 2.53-2.65 (m, 4H), 3.20-3.33 (m, 6H), 3.48-3.58 (m, 2H), 3.66 (s, 2H), 6.86-6.91 (m, 1H), 7.21-7.32 (m, 7H), 7.32-7.39 (m, 3H), 7.56-7.60 (m, 1H). Found: C, 57.66; H, 5.21; N, 5.88. C₃₁H₃₄N₃O₂Cl x 1.6 CF₃CO₂H x 0.8 H₂O has C, 57.62; H, 5.26; N, 5.89%.

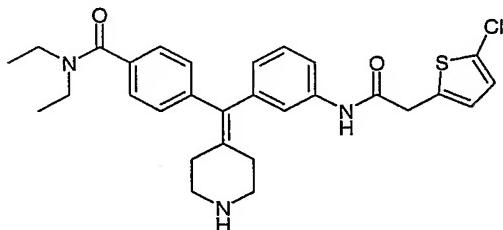
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COMPOUND 23: N,N-diethyl-4-[(3-{[(5-methylthien-2-yl)acetyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide

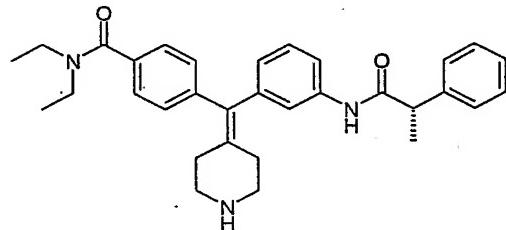


Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.140 g, 0.302 mmol) and (5-methylthien-2-yl)acetic acid (71 mg, 0.46 mmol) provided COMPOUND 23 (121 mg, 65%) as a colourless solid. Purity (HPLC): >99% (215 nm), >99% (254 nm), >99% (280 nm); ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.83 Hz, 3H), 1.23 (br t, J = 6.83 Hz, 3H), 2.41 (d, J = 0.78 Hz, 3H), 2.55-2.64 (m, 4H), 3.20-3.34 (m, 6H), 3.48-3.58 (m, 2H), 3.76 (s, 2H), 6.59-6.62 (m, 1H), 6.71-6.75 (m, 1H), 6.86-6.91 (m, 1H), 7.26 (d, J = 8.40 Hz, 2H), 7.27-7.33 (m, 2H), 7.35 (d, J = 8.40 Hz, 2H), 7.56-7.59 (m, 1H). Found: C, 59.92; H, 5.74; N, 6.46. C₃₀H₃₅N₃O₂S x 1.2 CF₃CO₂H x 0.6 H₂O has C, 59.93; H, 5.81; N, 6.47%.

COMPOUND 24: 4-[(3-{[(5-chlorothien-2-yl)acetyl]amino}phenyl)(piperidin-4-

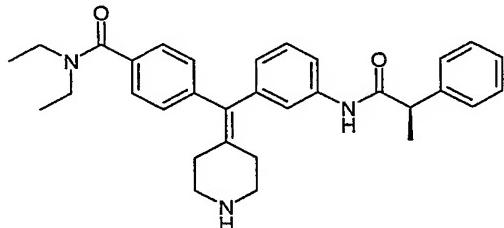
ylidene)methyl]-N,N-diethylbenzamide

Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.164 g, 0.354 mmol) and 5-chlorothiophene-2-acetic acid (86 mg, 0.53 mmol) provided
 5 COMPOUND 24 (168 mg, 75%) as a colourless solid. Purity (HPLC): >99% (215 nm), >99% (254 nm), >99% (280 nm); ^1H NMR (400 MHz, CD₃OD) δ 1.11 (br t, J = 7.03 Hz, 3H), 1.23 (br t, J = 7.03 Hz, 3H), 2.54-2.65 (m, 4H), 3.20-3.34 (m, 6H), 3.48-3.58 (m, 2H), 3.80 (d, J = 0.78 Hz, 2H), 6.77-6.79 (m, 1H), 6.82 (d, J = 3.71 Hz, 1H), 6.88-6.92 (m, 1H), 7.26 (d, J = 8.40 Hz, 2H), 7.28-7.34 (m, 2H), 7.36 (d, J = 8.20 Hz, 2H), 7.57-7.60 (m, 1H). Found: C, 54.64; H, 4.85; N, 5.92. C₂₉H₃₂N₃O₂SCl x 1.6 CF₃CO₂H x 0.2 H₂O has C, 54.51; H, 4.84; N, 5.93%.

COMPOUND 25: N,N-diethyl-4-[(3-{[(2S)-2-phenylpropanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide

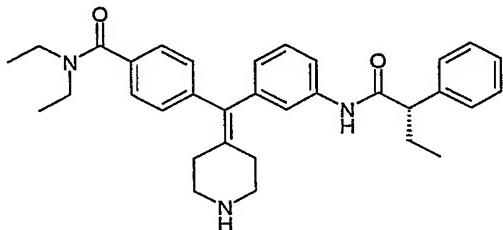
15 Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.152 g, 0.328 mmol) and (S)-(+)2-phenylpropionic acid (74 mg, 0.49 mmol) provided COMPOUND 25 (129 mg, 65%) as a colourless solid. Purity (HPLC): >99% (215 nm), >99% (254 nm), >99% (280 nm); Optical purity (Chiral HPLC): > 99%; ^1H NMR (400 MHz, CD₃OD) δ 1.11 (br t, J = 6.64 Hz, 3H), 1.22 (br t, J = 7.23 Hz, 3H), 1.49 (d, J = 7.03 Hz, 3H), 2.54-2.63 (m, 4H), 3.19-3.35 (m, 6H), 3.48-3.58 (m, 2H), 3.78 (q, J = 7.03 Hz, 1H), 6.84-6.89 (m, 1H), 7.20-7.28 (m, 4H), 7.27-7.40 (m, 7H), 7.53-7.56 (m, 1H). Found: C, 61.47; H, 5.79; N, 6.08. C₃₂H₃₇N₃O₂ x 1.6 CF₃CO₂H x 0.5 H₂O has C, 61.53; H, 5.81; N, 6.12%.

COMPOUND 26: *N,N*-diethyl-4-[(3-{[(2R)-2-phenylpropanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide



- 5 Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.309 g, 0.666 mmol) and (R)-(-)-2-phenylpropionic acid (0.150 g, 0.999 mmol) provided COMPOUND 26 (175 mg, 43%) as a colourless solid. Purity (HPLC): >99% (215 nm), >99% (254 nm), >99% (280 nm); Optical purity (Chiral HPLC): > 99%; ^1H NMR (400 MHz, CD₃OD) δ 1.11 (br t, J = 7.42 Hz, 3H), 1.23 (br t, J = 6.83 Hz, 3H), 1.48 (d, J = 7.03 Hz, 3H), 2.55-2.62 (m, 4H), 3.21-3.32 (m, 6H), 3.49-3.57 (m, 2H), 3.78 (q, J = 7.03 Hz, 1H), 6.86 (dt, J = 7.23, 1.37 Hz, 1H), 7.20-7.27 (m, 3H), 7.27-7.40 (m, 8H), 7.53-7.55 (m, 1H). Found: C, 60.28; H, 5.79; N, 6.16. C₃₂H₃₇N₃O₂ x 1.7 CF₃CO₂H x 0.9 H₂O has C, 60.25; H, 5.78; N, 5.95%.
- 10

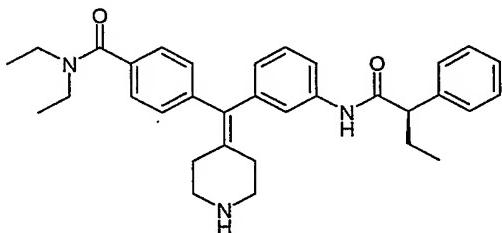
- 15 **COMPOUND 27: *N,N*-diethyl-4-[(3-{[(2S)-2-phenylbutanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide**



- Using the same method as for COMPOUND 20 and using INTERMEDIATE 6 (0.346 g, 0.746 mmol) and (S)-(+)-2-phenylbutyric acid (0.184 g, 1.12 mmol) provided
20 COMPOUND 27 (148 mg, 32%) as a colourless solid. Purity (HPLC): >99%; Optical purity (Chiral HPLC): > 99%; ^1H NMR (400 MHz, CD₃OD) δ 0.92 (t, J = 7.42 Hz, 3H), 1.11 (br t, J = 7.03 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 1.72-1.85 (m, 1H), 2.05-2.17 (m, 1H), 2.54-2.62 (m, 4H), 3.21-3.32 (m, 6H), 3.46-3.57 (m, 3H), 6.85-

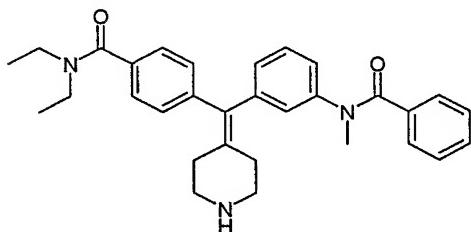
6.89 (m, 1H), 7.20-7.40 (m, 11H), 7.52-7.55 (m, 1H). Found: C, 51.82; H, 4.66; N, 4.75. $C_{33}H_{39}N_3O_2 \times 3.6 CF_3CO_2H \times 0.6 H_2O$ has C, 51.86; H, 4.74; N, 4.51%.

5 **COMPOUND 28: *N,N*-diethyl-4-[(3-{[(2R)-2-phenylbutanoyl]amino}phenyl)(piperidin-4-ylidene)methyl]benzamide**



Using the same method as for COMPOUND 19 and using INTERMEDIATE 6 (0.309 g, 0.666 mmol) and (R)-(-)-2-phenylbutyric acid (0.354 g, 0.932 mmol) provided COMPOUND 28 (189 mg, 45%) as a colourless solid. Purity (HPLC): >99%; Optical purity (Chiral HPLC): > 99%; 1H NMR (400 MHz, CD₃OD) δ 0.92 (t, J = 7.42 Hz, 3H), 1.11 (br t, J = 7.62 Hz, 3H), 1.23 (br t, J = 6.64 Hz, 3H), 1.72-1.84 (m, 1H), 2.06-2.16 (m, 1H), 2.55-2.62 (m, 4H), 3.21-3.32 (m, 6H), 3.47-3.57 (m, 3H), 6.87 (dt, J = 7.42, 1.37 Hz, 1H), 7.22-7.33 (m, 7H), 7.34 (d, J = 8.40 Hz, 2H), 7.36-7.40 (m, 2H), 7.53-7.55 (m, 1H). Found: C, 64.16; H, 6.79; N, 6.26. $C_{33}H_{39}N_3O_2 \times 1.0 CF_3CO_2H \times 1.8 H_2O$ has C, 64.07; H, 6.70; N, 6.40%.

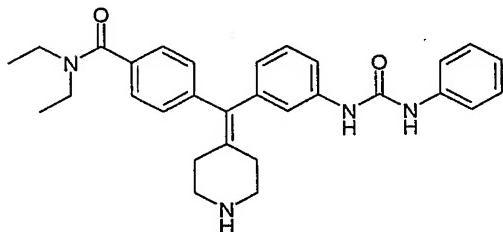
COMPOUND 29: 4-[(3-[benzoyl(methyl)amino]phenyl)(piperidin-4-ylidene)methyl]-*N,N*-diethylbenzamide



20 To a solution of INTERMEDIATE 6 (0.153 g, 0.330 mmol) in dichloromethane (10 mL) was added benzoyl chloride (42 μ L, 0.36 mmol) and triethylamine (0.15 mL, 1.1 mmol). The reaction was stirred for 24 h at room temperature under an atmosphere of nitrogen. The mixture was washed with saturated aqueous sodium bicarbonate and the aqueous layer extracted with two portions of dichloromethane. The combined

organic extracts were dried (Na_2SO_4), filtered and concentrated. The crude material (0.186 g, 0.330 mmol) was dissolved in THF (5 mL). To the solution was added sodium hydride (60% dispersion in mineral oil) (12 mg, 0.30 mmol). The reaction was stirred for 1 hour under N_2 , after which methyl iodide (20 μL , 0.32 mmol) was added. The reaction was stirred overnight at room temperature. Saturated aqueous ammonium chloride was added and the reaction mixture was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na_2SO_4), filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (0.5 mL) was added. After 2 hours, the solution was concentrated and the residue was purified by reverse phase chromatography, eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 29 (56 mg, 74%) as an off-white solid. Purity (HPLC) >99%; ^1H NMR (400 MHz, CD_3OD) δ 1.12 (t, $J = 6.64$ Hz, 3H), 1.24 (t, $J = 6.64$ Hz, 3H), 2.00-2.09 (m, 2H), 2.44-2.52 (m, 2H), 2.98-3.06 (m, 2H), 3.14-3.21 (m, 2H), 3.23-3.34 (m, 2H), 3.47 (s, 3H), 3.49-3.58 (m, 2H), 6.82 (br s, 1H), 6.91 (m, 1H), 7.06 (d, $J = 8.01$ Hz, 2H), 7.19-7.38 (m, 9H). Found: C, 52.24; H, 5.27; N, 4.96. $\text{C}_{31}\text{H}_{35}\text{N}_3\text{O}_2 \times 2.6 \text{ CF}_3\text{CO}_2\text{H} \times 3.0 \text{ H}_2\text{O}$ has C, 52.25; H, 5.28; N, 5.05%.

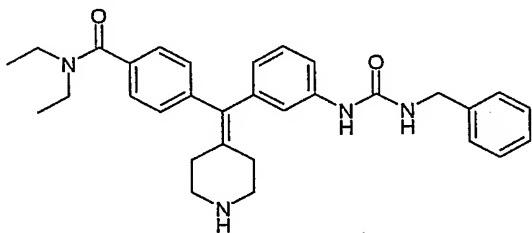
20 **COMPOUND 30: 4-[{3-[{(anilinocarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide}**



To a solution of the INTERMEDIATE 6 (0.127 g, 0.274 mmol) in DMF was added phenylisocyanate (38 μL , 0.36 mmol). The reaction was stirred overnight at room temperature under nitrogen. The reaction mixture was concentrated and the residue was diluted with dichloromethane. The solution was washed with one portion of saturated aqueous ammonium chloride and the organic phase was dried (Na_2SO_4), filtered and concentrated. The residue was dissolved in dichloromethane (5 mL) and

trifluoroacetic acid (1 mL) was added. After 4 hours, the solution was concentrated and purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as its TFA salt and was lyophilized to give COMPOUND 30 (123 mg, 75%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.83 Hz, 3H), 1.23 (br t, J = 6.83 Hz, 3H), 2.55-2.67 (m, 4H), 3.22-3.35 (m, 6H), 3.47-4.59 (m, 2H), 6.78-6.82 (m, 1H), 6.99-7.04 (m, 1H), 7.15 (ddd, J = 8.20, 2.15, 0.98 Hz, 1H), 7.23-7.31 (m, 5H), 7.36 (d, J = 8.40 Hz, 2H), 7.38-7.42 (m, 2H), 7.49-7.51 (m, 1H). Found: C, 60.78; H, 5.63; N, 8.66. C₃₀H₃₄N₄O₂ x 1.4 CF₃CO₂H x 0.3 H₂O has C, 60.83; H, 5.60; N, 8.65%.

COMPOUND 31: 4-[(3-{[(benzylamino)carbonyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide

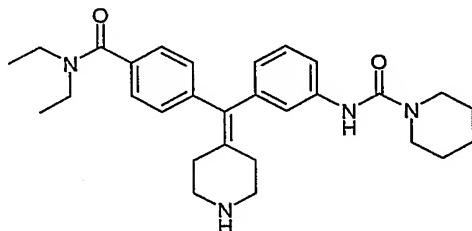


To a solution of INTERMEDIATE 6 (0.350 g, 0.755 mmol) in 1,2-dichloroethane was added benzyl isocyanate (0.14 mL, 1.13 mmol). The reaction was stirred overnight at room temperature under nitrogen. Trifluoroacetic acid (2 mL) was then added. After 4 hours, saturated aqueous sodium bicarbonate was added until bubbling ceased. The layers were separated and the aqueous layer was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na₂SO₄), filtered and concentrated. The residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as its TFA salt and was lyophilized to give COMPOUND 31 (201 mg, 44%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.03 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 2.55-2.65 (m, 4H), 3.21-3.28 (m, 4H), 3.28-3.33 (m, 2H), 3.49-3.59 (m, 2H), 4.37 (s, 2H), 6.75 (ddd, J = 2.73, 1.56, 1.17 Hz, 1H), 7.10 (ddd, J = 8.01, 2.15, 0.98 Hz, 1H), 7.19-7.25 (m, 3H), 7.27 (d, J = 8.40 Hz, 2H), 7.29-7.33 (m, 3H), 7.35 (d, J = 8.40 Hz,

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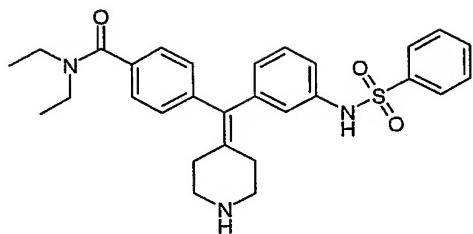
2H), 7.42-7.45 (m, 1H). Found: C, 60.76; H, 5.81; N, 8.35. $C_{31}H_{36}N_4O_2 \times 1.4$ CF₃CO₂H × 0.7 H₂O has C, 60.69; H, 5.85; N, 8.38%.

COMPOUND 32: N-[3-[{4-[(diethylamino)carbonyl]phenyl}(piperidin-4-ylidene)methyl]phenyl]piperidine-1-carboxamide



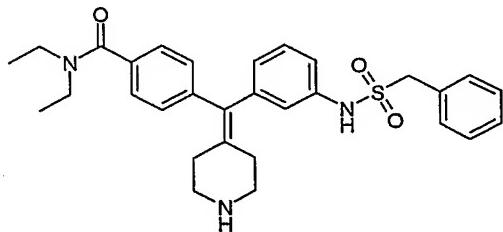
To a solution of INTERMEDIATE 6 (0.113 g, 0.244 mmol) in pyridine (8 mL) was added 1-piperidincarbonyl chloride (34 µL, 0.27 mmol). The reaction was stirred for 18 hours at room temperature under nitrogen. The mixture was concentrated and the residue was diluted with dichloromethane. The solution was washed with 1M HCl (1x) and the organic layer was dried (Na₂SO₄), filtered and concentrated. The residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1 mL) was added. After 2 hours, the reaction mixture was concentrated and the residue was purified by reverse phase chromatography (gradient 10-45% acetonitrile in water containing 0.1% trifluoroacetic acid). The product was obtained as its TFA salt and was lyophilized to give COMPOUND 32 (36 mg, 25%) as a colourless solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 8.81 Hz, 3H), 1.23 (br t, J = 7.81 Hz, 3H), 1.53-1.63 (m, 4H), 1.63-1.72 (m, 2H), 2.52-2.67 (m, 4H), 3.20-3.33 (m, 6H), 3.47-3.50 (m, 4H), 3.54-3.58 (m, 2H), 6.76-6.80 (m, 1H), 7.15-7.20 (m, 1H), 7.20-7.25 (m, 1H), 7.27 (d, J = 8.40 Hz, 2H), 7.29-7.31 (m, 1H), 7.35 (d, J = 8.40 Hz, 2H). Found: C, 56.68; H, 6.02; N, 7.93. $C_{29}H_{38}N_4O_2 \times 1.8$ CF₃CO₂H × 0.6 H₂O has C, 56.69; H, 5.98; N, 8.11%.

COMPOUND 33: N,N-diethyl-4-[{3-[(phenylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide



A solution of INTERMEDIATE 6 (0.119 g, 0.257 mmol), benzenesulfonyl chloride (36 μ L, 0.28 mmol) and triethylamine (0.11 mL, 0.77 mmol) in dichloromethane (10 mL) was stirred overnight at room temperature under nitrogen. Trifluoroacetic acid 5 (1 mL) was added and the reaction was stirred for an additional 2 hours. Saturated aqueous sodium bicarbonate was added until bubbling ceased. The layers were separated and the aqueous layer was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na_2SO_4), filtered and concentrated. The residue was purified by reverse phase chromatography, eluting 10% to 45% 10 acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 33 (63 mg, 40%) as a colourless solid. Purity (HPLC) >99%; ^1H NMR (400 MHz, CD_3OD) δ 1.09 (t, J = 6.44 Hz, 3H), 1.20 (t, J = 7.23 Hz, 3H), 2.40-2.46 (m, 2H), 2.49-2.56 (m, 2H), 3.13-3.24 (m, 6H), 3.46-3.55 (m, 2H), 6.82 (ddd, J = 7.62, 1.56, 0.98 Hz, 1H), 6.89 (m, 1H), 6.94 (m, 1H), 7.11-7.18 (m, 3H), 7.32 (d, J = 8.40 Hz, 2H), 7.38-7.45 (m, 2H), 15 7.52 (m, 1H), 7.59-7.64 (m, 2H). Found: C, 44.05; H, 4.28; N, 4.04. $\text{C}_{29}\text{H}_{33}\text{N}_3\text{O}_3\text{S} \times 4.0 \text{ CF}_3\text{CO}_2\text{H} \times 2.7 \text{ H}_2\text{O}$ has C, 44.07; H, 4.24; N, 4.17%.

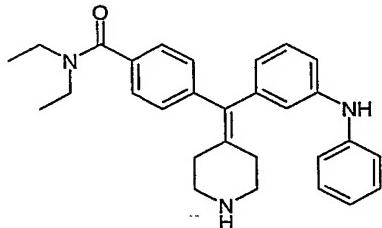
COMPOUND 34: 4-[{3-[(benzylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



Using the same method as for COMPOUND 33 and using INTERMEDIATE 6 (0.104 g, 0.225 mmol) and α -toluenesulfonyl chloride (47 mg, 0.25 mmol) provided COMPOUND 34 (64 mg, 45%) as a colourless solid. Purity (HPLC): >99%; ^1H

NMR (400 MHz, CD₃OD) δ 1.09 (t, J = 6.83 Hz, 3H), 1.22 (t, J = 7.03 Hz, 3H), 2.55-2.62 (m, 4H), 3.21-3.32 (m, 6 H), 3.48-3.58 (m, 2 H), 4.34 (s, 2H), 6.88 (m, 1H), 6.96 (m, 1H), 7.04 (dd, J = 2.15, 0.98 Hz, 1H), 7.19-7.35 (m, 8H), 7.38 (d, J = 8.20 Hz, 2H). Found: C, 48.57; H, 5.04; N, 4.27. C₃₀H₃₅N₃O₃S x 2.8 CF₃CO₂H x 2.5 H₂O has C, 48.48; H, 4.89; N, 4.76%.

COMPOUND 35: 4-[(3-anilinophenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide

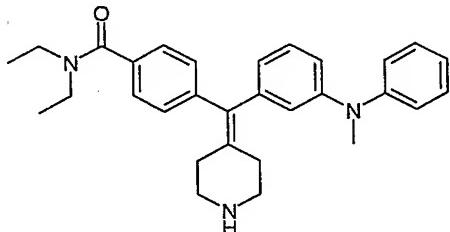


A SmithProcess vial was charged with INTERMEDIATE 6 (0.205 g, 0.442 mmol), bromobenzene (61 μL, 0.58 mmol), Pd₂(dba)₃ (16 mg, 0.017 mmol), sodium *tert*-butoxide (60 mg, 0.63 mmol), BINAP (22 mg, 0.035 mmol) and toluene (2.5 mL) and the cap was tightened thoroughly. The vessel was exposed to microwave irradiation in a SmithSynthesizer for 5 min at 110 °C. The reaction tube was cooled to room temperature and the mixture was absorbed onto silica gel. Flash chromatography (eluting 20% to 50% ethyl acetate in hexanes) afforded the BOC-protected compound as a yellow oil. The oil was dissolved in dichloromethane (15 mL) and trifluoroacetic acid was added (1 mL). After 3 hours of stirring, the reaction was neutralized with 2M NaOH. The phases were separated and the aqueous phase was extracted with two portions of dichloromethane. The combined organic extracts were dried (Na₂SO₄), filtered and concentrated to afford the pure product as a yellow oil. The product was dissolved in a 1:5 mixture of dichloromethane/ether (12 mL) and 2.2 mL of 1M HCl in ether was added under an atmosphere of nitrogen. Concentration of the solution provided COMPOUND 35 (208 mg, 92%) as its HCl salt. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.24 (br t, J = 6.64 Hz, 3H), 2.55-2.62 (m, 2H), 2.63-2.70 (m, 2H), 3.21-3.35 (m, 6H), 3.50-3.58 (m, 2H), 6.61-6.67 (m, 1H), 6.81-6.87 (m, 2H), 6.95-7.04 (m, 3H), 7.15-7.22 (m, 3H), 7.27 (d, J = 8.40 Hz, 2H), 7.36 (d, J = 8.40 Hz, 2H). Found: C,

63

70.20; H, 7.26; N, 8.33. $C_{29}H_{33}N_3O \times 1.2\text{ HCl} \times 0.7\text{ H}_2O$ has C, 70.23; H, 7.24; N, 8.47%.

5 **COMPOUND 36: *N,N*-diethyl-4-[{3-[methyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide**

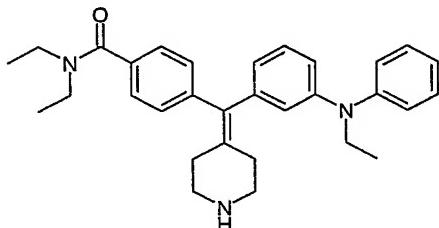


A SmithProcess vial was charged with INTERMEDIATE 6 (0.205 g, 0.442 mmol), bromobenzene (61 μL , 0.58 mmol), $\text{Pd}_2(\text{dba})_3$ (16 mg, 0.017 mmol), sodium *tert*-butoxide (60 mg, 0.63 mmol), BINAP (22 mg, 0.035 mmol) and toluene (2.5 mL) and the cap was tightened thoroughly. The vessel was exposed to microwave irradiation in a SmithSynthesizer for 5 min at 110 °C. The reaction tube was cooled to room temperature and the mixture was absorbed onto silica gel. Flash chromatography (eluting 0% to 50% ethyl acetate in hexanes) afforded the N-arylated intermediate as a yellow oil (0.238 g, 0.441 mmol). The oil was dissolved in dry DMF (10 mL) and NaH (60% dispersion in mineral oil) (42 mg, 1.0 mmol) was added. The reaction was stirred for 1 h, after which iodomethane (69 μL , 1.1 mmol) was added. After 4 h, saturated aqueous NH_4Cl was added. The mixture was diluted with dichloromethane and washed with saturated aqueous NH_4Cl . The aqueous layer was extracted with two portions of dichloromethane and the combined organic extracts were dried (Na_2SO_4), filtered and concentrated. The residue was then dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1 mL) was added. After 18 h, the solution was concentrated and purified by reverse phase chromatography, eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 36 (210 mg, 70% yield) as a yellow solid. Purity (HPLC) >99%; ^1H NMR (400 MHz, CD_3OD) δ 1.12 (t, $J = 7.23\text{ Hz}$, 3H), 1.23 (t, $J = 7.23\text{ Hz}$, 3H), 2.51-2.57 (m, 2H), 2.57-2.63 (m, 2H), 3.16-3.23 (m, 4H), 3.25 (s, 3H), 3.26-3.34 (m, 2H), 3.48-3.58 (m, 2H), 6.64-6.69 (m, 1H), 6.69-6.72 (m, 1H), 6.89

64

(ddd, $J = 8.40, 2.54, 0.98$ Hz, 1H), 6.95-7.00 (m, 1H), 7.00-7.03 (m, 2H), 7.19-7.28 (m, 5H), 7.34 (d, $J = 8.40$ Hz, 2H). Found: C, 63.98; H, 6.26; N, 6.79. $C_{30}H_{35}N_3O \times 1.3 CF_3CO_2H \times 0.6 H_2O$ has C, 63.91; H, 6.17; N, 6.86%.

5 COMPOUND 37: *N,N*-diethyl-4-[{3-[ethyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide



A SmithProcess vial was charged with INTERMEDIATE 6 (0.400 g, 0.863 mmol), bromobenzene (118 μ L, 1.12 mmol), Pd₂(dba)₃ (32 mg, 0.035 mmol), sodium *tert*-butoxide (116 mg, 0.121 mmol), BINAP (43 mg, 0.069 mmol) and toluene (3.0 mL) and the cap was tightened thoroughly. The vessel was exposed to microwave irradiation in a SmithSynthesizer for 5 min at 110 °C. The reaction tube was cooled to room temperature and the mixture was absorbed onto silica gel. Flash chromatography (eluting 0% to 50% ethyl acetate in hexanes) afforded the N-arylated intermediate as a yellow oil (0.424 g, 0.786 mmol). The oil was dissolved in dry DMF (20 mL) and NaH (60% dispersion in mineral oil) (75 mg, 1.9 mmol) was added. The reaction was stirred for 30 minutes, after which ethyl iodide (0.16 mL, 2.0 mmol) was added. After 4 h, the reaction mixture was diluted with ethyl acetate and washed with saturated aqueous NH₄Cl. The organic layer was dried (Na₂SO₄), filtered and concentrated. The residue was then dissolved in dichloromethane (10 mL) and trifluoroacetic acid (2 mL) was added. After 18 h, the solution was concentrated and purified by reverse phase chromatography eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as the trifluoroacetic acid salt and was lyophilized to give COMPOUND 37 (321 mg, 70% yield) as a yellow solid. Purity (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, $J = 6.64$ Hz, 3H), 1.15 (t, $J = 7.03$ Hz, 3H), 1.24 (br t, $J = 7.42$ Hz, 3H), 2.52-2.57 (m, 2H), 2.58-2.63 (m, 2H), 3.16-3.24 (m, 4H), 3.24-3.34 (m, 2H), 3.49-3.59 (m, 2H), 3.76 (q, $J = 7.03$ Hz, 2H), 6.61-6.66 (m, 2H), 6.86 (ddd, $J = 8.40, 2.54,$

65

0.98 Hz, 1H), 6.95-6.98 (m, 1H), 6.99-7.03 (m, 2H), 7.17-7.28 (m, 5H), 7.34 (d, J = 8.40 Hz, 2H). Found: C, 63.42; H, 6.03; N, 6.53. $C_{31}H_{37}N_3O \times 1.5 CF_3CO_2H \times 0.3 H_2O$ has C, 63.40; H, 6.12; N, 6.52%.

5 **INTERMEDIATE 7: *tert*-butyl 4-[{4-[(diethylamino)carbonyl]phenyl}(3-hydroxyphenyl)methylene]piperidine-1-carboxylate**

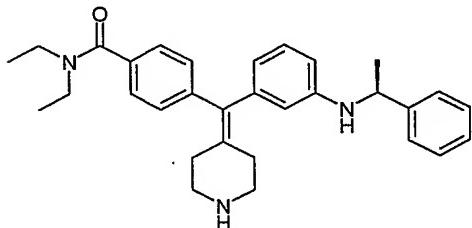
A solution of INTERMEDIATE 5 (4.08 g, 9.04 mmol), 3-hydroxyphenylboronic acid (1.97 g, 14.3 mmol) and aqueous 2N sodium carbonate (11.3 mL, 22.6 mmol) in a 1:1
10 mixture of toluene/ethanol (200 mL) was degassed for 20 minutes. Palladium tetrakis(triphenylphosphine) (1.05 g, 0.909 mmol) was added and the reaction mixture was purged with nitrogen and heated to 90 °C. After 5 h, the reaction was cooled to room temperature and saturated aqueous ammonium chloride was added. The mixture was extracted with two portions of ethyl acetate and the combined organic
15 extracts were dried (Na_2SO_4), filtered and concentrated. The residue was purified by flash chromatography eluting with 0% to 100% ethyl acetate in hexanes to yield
INTERMEDIATE 7 as a white solid (4.24 g, 100%). 1H NMR (400MHz, $CDCl_3$) δ 1.10 (t, J = 7.42 Hz, 3H), 1.20 (t, J = 7.03 Hz, 3H), 1.42 (s, 9H), 2.25-2.33 (m, 4H),
3.23-3.31 (m, 2H), 3.39-3.46 (m, 4H), 3.46-3.54 (m, 2H), 6.51 (dd, J = 2.15, 1.56 Hz,
20 1H), 6.57 (ddd, J = 7.62, 1.56, 0.98 Hz, 1H), 6.62 (ddd, J = 8.20, 2.54, 0.98 Hz, 1H),
7.06-7.12 (m, 1H), 7.19 (d, J = 8.40 Hz, 2H), 7.29 (d, J = 8.40 Hz, 2H).

INTERMEDIATE 8: *tert*-butyl 4-[{4-[(diethylamino)carbonyl]phenyl}(3-[(trifluoromethyl)sulfonyl]oxy)phenyl)methylene]piperidine-1-carboxylate

25 To a solution of INTERMEDIATE 6 (1.95 g, 4.20 mmol) and triethylamine (3.00 mL, 21.5 mmol) in dichloromethane (15 mL) was added trifluoromethanesulfonic anhydride (0.95 mL, 5.65 mmol) dropwise at 0 °C. The reaction was allowed to warm to room temperature and was stirred for 15 h. The solution was diluted with dichloromethane and was washed with two portions of saturated aqueous ammonium
30 chloride, then dried (Na_2SO_4), filtered and concentrated. The residue was purified by flash chromatography (eluting 20% to 50% ethyl acetate in hexanes) to yield
INTERMEDIATE 8 (2.00 g, 80%) as a yellow foam. 1H NMR (400MHz, $CDCl_3$) δ

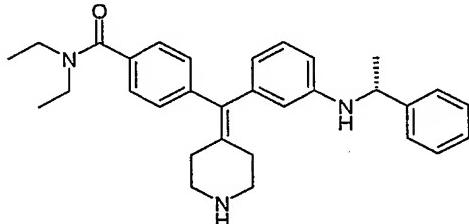
1.08-1.18 (m, 3H), 1.20-1.29 (m, 3H), 1.47 (s, 9H), 2.29-2.38 (m, 4H), 3.23-3.34 (m, 2H), 3.44-3.50 (m, 4H), 3.50-3.60 (m, 2H), 7.02-7.05 (m, 1H), 7.09-7.18 (m, 4H), 7.34 (d, J = 8.01 Hz, 2H), 7.36-7.41 (m, 1H).

5 **COMPOUND 38: N,N-diethyl-4-[(3-[(1*S*)-1-phenylethyl]amino)phenyl](piperidin-4-ylidene)methyl]benzamide**



Into an oven-dried test tube were weighed Pd₂(dba)₃ (7.6 mg, 0.0083 mmol), BINAP (10.7 mg, 0.0172 mmol) and sodium *tert*-butoxide (24.9 mg, 0.260 mmol). The solid reagents were suspended in degassed toluene (3 mL). The test tube was sealed with a rubber septum and purged with nitrogen. (S)-(-)-1-phenylethylamine (30 µL, 0.23 mmol) was added to the mixture by syringe. The reaction was heated at 80 °C for 5 minutes. A solution of INTERMEDIATE 8 (97 mg, 0.16 mmol) was added dropwise by syringe over a 15 minute period. The reaction was stirred for 8 hours, then cooled to room temperature. The mixture was purified by flash chromatography eluting with 20% to 50% ethyl acetate in hexanes to give 66 mg of a yellow oil. The oil was dissolved in dichloromethane (8 mL) and trifluoroacetic acid (0.5 mL) was added. After 4 hours, the reaction mixture was concentrated and the residue was purified by reverse phase chromatography eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 38 (88 mg, 78%) as a colourless solid. Purity (HPLC) >99%; Optical purity (Chiral HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.23 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 1.51 (d, J = 6.83 Hz, 3H), 2.29-2.36 (m, 2H), 2.46-2.52 (m, 2H), 2.90-3.09 (m, 2H), 3.09-3.22 (m, 2H), 3.24-3.36 (m, 2H), 3.48-3.59 (m, 2H), 4.49 (q, J = 6.83 Hz, 1H), 6.31-6.34 (m, 1H), 6.36-6.41 (m, 1H), 6.59-6.65 (m, 1H), 7.04 (t, J = 8.01 Hz, 1H), 7.13 (d, J = 8.40 Hz, 2H), 7.17-7.23 (m, 1H), 7.25-7.34 (m, 6H). Found: C, 58.75; H, 5.56; N, 5.83. C₃₁H₃₇N₃O x 2.2 CF₃CO₂H x 0.3 H₂O has C, 58.74; H, 5.54; N, 5.80%.

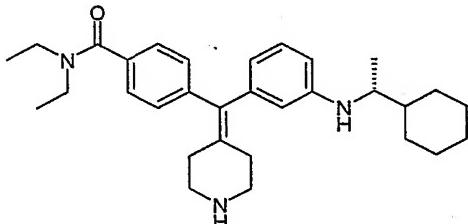
COMPOUND 39: *N,N*-diethyl-4-[(3-[(1*R*)-1-phenylethyl]amino)phenyl](piperidin-4-ylidene)methyl]benzamide



5 Using the same method as for COMPOUND 38 and using INTERMEDIATE 8 (0.398 g, 0.667 mmol) and (R)-(+)2-phenylethylamine (0.14 mL, 1.0 mmol) provided COMPOUND 39 (101 mg, 22%) as a colourless solid. Purity (HPLC): >99%; Optical purity (Chiral HPLC): > 99%; ^1H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.23 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 1.51 (d, J = 6.83 Hz, 3H), 2.29-2.36 (m, 2H), 2.46-10 2.52 (m, 2H), 2.90-3.09 (m, 2H), 3.09-3.22 (m, 2H), 3.24-3.36 (m, 2H), 3.48-3.59 (m, 2H), 4.49 (q, J = 6.83 Hz, 1H), 6.31-6.34 (m, 1H), 6.36-6.41 (m, 1H), 6.59-6.65 (m, 1H), 7.04 (t, J = 8.01 Hz, 1H), 7.13 (d, J = 8.40 Hz, 2H), 7.17-7.23 (m, 1H), 7.25-7.34 (m, 6H). Found: C, 49.48; H, 4.89; N, 4.58. C₃₁H₃₇N₃O x 3.7 CF₃CO₂H x 2.4 H₂O has C, 49.45; H, 4.92; N, 4.50%.

15

COMPOUND 40: 4-[(3-[(1*R*)-1-cyclohexylethyl]amino)phenyl](piperidin-4-ylidene)methyl]-*N,N*-diethylbenzamide



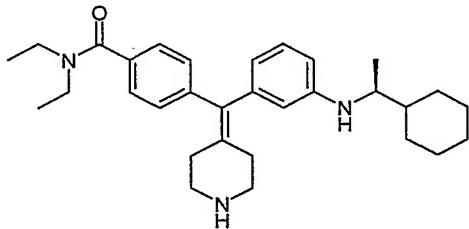
In a 2-necked flask equipped with a reflux condenser were added Pd₂(dba)₃ (30 mg, 0.033 mmol), BINAP (42 mg, 0.067 mmol) and sodium *tert*-butoxide (96 mg, 1.00 mmol). The solid materials were suspended in degassed toluene (10 mL). (R)-(-)-1-cyclohexylethylamine (0.14 mL, 0.94 mmol) was added by syringe. The reaction was heated at 80 °C for 5 minutes. A solution of INTERMEDIATE 8 (0.400 g, 0.670 mmol) was added dropwise by syringe over a 20 minute period. The reaction was 20 stirred for 15 h and then cooled to room temperature. The mixture was purified by 25

flash chromatography eluting with 0% to 50% ethyl acetate in hexanes to give 111 mg of a yellow oil. The oil was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (1.0 mL) was added. After 4 hours, the reaction mixture was concentrated and the residue was purified by reverse phase chromatography eluting 10% to 45%

5 acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 38 (100 mg, 21%) as a colourless solid. Purity (HPLC) >99%; Optical purity (Chiral HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.08-1.28 (m, 11H), 1.17 (d, J = 6.83 Hz, 3H), 1.52-1.85 (m, 6H), 2.57-2.65 (m, 4H), 3.23-3.34 (m, 6H), 3.41-3.49 (m, 1H), 3.49-3.59 (m, 2H),

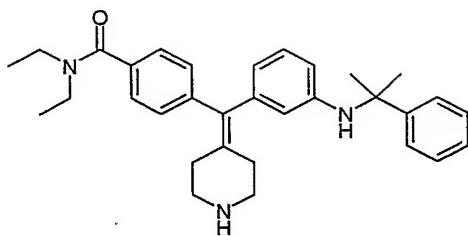
10 7.05-7.08 (m, 1H), 7.19-7.24 (m, 2H), 7.28 (d, J = 8.40 Hz, 2H), 7.38 (d, J = 8.40 Hz, 2H), 7.47-7.53 (m, 1H). Found: C, 58.42; H, 6.22; 5.78. C₃₁H₄₃N₃O x 2.2 CF₃CO₂H x 0.2 H₂O has C, 58.39; H, 6.31; N, 5.77%.

15 **COMPOUND 41: 4-[(3-{[(1S)-1-cyclohexylethyl]amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide**



Using the same method as for COMPOUND 40 and using INTERMEDIATE 8 (0.400 g, 0.670 mmol) and (R)-(-)-1-cyclohexylethylamine (0.14 mL, 0.94 mmol) provided COMPOUND 41 (30 mg, 6%) as a colourless solid. Purity (HPLC) >97% (215 nm), >97% (254 nm), >99% (280 nm); Optical purity (Chiral HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.08-1.28 (m, 11H), 1.17 (d, J = 6.83 Hz, 3H), 1.52-1.85 (m, 6H), 2.57-2.65 (m, 4H), 3.23-3.34 (m, 6H), 3.41-3.49 (m, 1H), 3.49-3.59 (m, 2H), 7.05-7.08 (m, 1H), 7.19-7.24 (m, 2H), 7.28 (d, J = 8.40 Hz, 2H), 7.38 (d, J = 8.40 Hz, 2H), 7.47-7.53 (m, 1H). Found: C, 59.88; H, 6.57; N, 5.98. C₃₁H₄₃N₃O x 1.9 CF₃CO₂H x 0.4 H₂O has C, 59.92; H, 6.60; N, 6.02%.

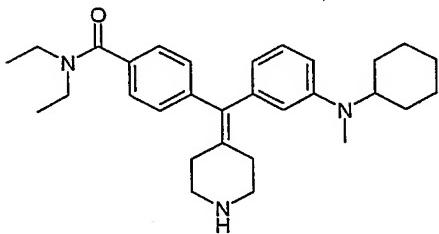
COMPOUND 42: N,N-diethyl-4-[(3-[(1-methyl-1-phenylethyl)amino]phenyl)(piperidin-4-ylidene)methyl]benzamide



In a 2-necked flask equipped with a reflux condenser were added INTERMEDIATE 8 (0.560 g, 0.939 mmol), Pd₂(dba)₃ (43 mg, 0.047 mmol), 2-(di-*t*-butylphosphino)biphenyl (28 mg, 0.094 mmol) and potassium phosphate (0.280 g, 1.32 mmol). The solid materials were suspended in degassed toluene (10 mL).

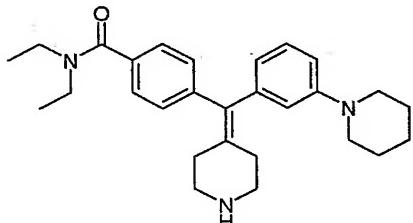
5 Cumylamine (0.152 g, 1.12 mmol) was added by syringe. The reaction was heated at 80 °C for 18 hours and then cooled to room temperature. The solution was diluted with ethyl acetate and washed with saturated aqueous sodium bicarbonate (1x) and brine (1x). The organic phase was dried (Na₂SO₄), filtered and concentrated. The 10 residue was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (2.0 mL) was added. After 4 hours, the reaction mixture was concentrated and the residue was purified by reverse phase chromatography eluting 10% to 45% acetonitrile in water containing 0.1% trifluoroacetic acid. The product was obtained as its TFA salt and was lyophilized to give COMPOUND 42 (69 mg, 10%) as a colourless solid. Purity 15 (HPLC) >99%; ¹H NMR (400 MHz, CD₃OD) δ 1.13 (br t, J = 7.23 Hz, 3H), 1.24 (br t, J = 6.64 Hz, 3H), 1.76 (s, 6H), 2.33-2.39 (m, 2H), 2.47-2.53 (m, 2H), 3.10-3.16 (m, 2H), 3.16-3.22 (m, 2H), 3.24-3.34 (m, 2H), 3.49-3.59 (m, 2H), 6.33-6.38 (m, 1H), 6.83 (dd, J = 7.81 Hz, 1.37 Hz, 1H), 6.96-7.02 (m, 1H), 7.08 (d, J = 8.40 Hz, 2H), 7.24-7.31 (m, 1H), 7.31-7.39 (m, 5H), 7.41-7.47 (m, 2H). Found: C, 55.34; H, 5.25; 20 N, 5.09. C₃₂H₃₉N₃O x 2.9 CF₃CO₂H x 0.4 H₂O has C, 55.40; H, 5.25; N, 5.13%.

COMPOUND 43: 4-[{3-[cyclohexyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide



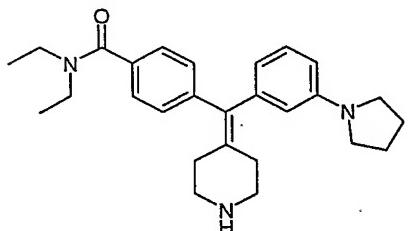
Using the same method as for COMPOUND 40 and using INTERMEDIATE 8 (0.400 g, 0.670 mmol) and *N*-methylcyclohexylamine (130 μ L, 0.996 mmol) provided COMPOUND 43 (37 mg, 8%) as a colourless solid. Purity (HPLC) >94% (215 nm), >95% (254 nm), >99% (280 nm); 1 H NMR (400 MHz, CD₃OD) δ 1.06-1.18 (m, 4H), 1.18-1.37 (m, 8H), 1.63-1.71 (m, 1H), 1.81-2.01 (m, 3H), 2.57-2.66 (m, 4H), 3.20-3.24 (s, 3H), 3.25-3.35 (m, 6H), 3.49-3.62 (m, 3H), 7.26-7.30 (m, 3H), 7.35-7.40 (m, 3H), 7.45 (ddd, J = 8.20, 2.34, 0.59 Hz, 1H), 7.56-7.63 (m, 1H). Found: C, 50.16; H, 5.26; N, 5.01. C₃₀H₄₁N₃O x 3.6 CF₃CO₂H x 1.1 H₂O has C, 50.20; H, 5.30; N, 4.72%.

10 **COMPOUND 44: *N,N*-diethyl-4-[piperidin-4-ylidene(3-piperidin-1-ylphenyl)methyl]benzamide**



Using the same method as for COMPOUND 40 and using INTERMEDIATE 8 (0.400 g, 0.670 mmol) and piperidine (100 μ L, 1.01 mmol) provided COMPOUND 44 (200 mg, 45%) as an orange solid. Purity (HPLC) >99%; 1 H NMR (400 MHz, CD₃OD) δ 1.13 (br t, J = 7.03 Hz, 3H), 1.23 (br t, J = 7.23 Hz, 3H), 1.73-1.81 (m, 2H), 1.95-2.04 (m, 4H), 2.55-2.64 (m, 4H), 3.24-3.33 (m, 6H), 3.49-3.61 (m, 6H), 7.28 (d, J = 8.40 Hz, 2H), 7.25-7.40 (m, 3H), 7.42-7.45 (m, 1H), 7.53-7.62 (m, 2H). Found: C, 50.78; H, 5.55; N, 5.14. C₂₈H₃₇N₃O x 2.9 CF₃CO₂H x 2.1 H₂O has C, 50.74; H, 5.56; N, 5.25%.

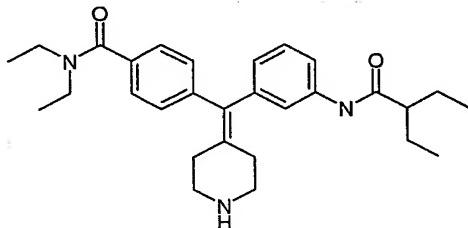
20 **COMPOUND 45: *N,N*-diethyl-4-[piperidin-4-ylidene(3-pyrrolidin-1-ylphenyl)methyl]benzamide**



Using the same method as for COMPOUND 40 and using INTERMEDIATE 8 (0.400 g, 0.670 mmol) and pyrrolidine (56 μ L, 1.0 mmol) provided COMPOUND 45 (174 mg, 40%) as a yellow solid. Purity (HPLC) >99%; 1 H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 7.62 Hz, 3H), 1.24 (br t, J = 7.62 Hz, 3H), 1.98-2.04 (m, 4H), 2.56-2.64 (m, 4H), 3.20-3.34 (m, 10H), 3.49-3.58 (m, 2H), 6.38-6.40 (m, 1H), 6.47-6.51 (m, 1H), 6.57 (ddd, J = 8.01, 2.34, 0.59 Hz, 1H), 7.14-7.20 (m, 1H), 7.28 (d, J = 8.40 Hz, 2H), 7.35 (d, J = 8.40 Hz, 2H). Found: C, 52.86; H, 5.45; N, 5.76. C₂₇H₃₅N₃O x 2.6 CF₃CO₂H x 1.0 H₂O has C, 52.83; H, 5.45; N, 5.74%.

10

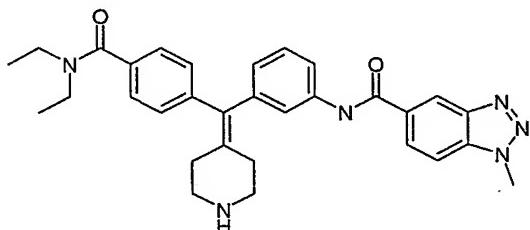
COMPOUND 46: N,N-diethyl-4-[[3-[(2-ethyl-1-oxobutyl)amino]phenyl]-4-piperidinylidenemethyl]-benzamide



15 To a solution of INTERMEDIATE 6 (225 mg, 0.485 mmol) and 2-ethylbutyryl chloride (98 mg, 0.728 mmol) in 1,2-dichloroethane (20 mL) was added triethylamine (0.203 mL, 1.456 mmol). The reaction was heated overnight at 50 °C, diluted with CH₂Cl₂ and washed with saturated aqueous NaHCO₃ (1x). The layers were separated and the aqueous layer was extracted with additional CH₂Cl₂ (1x). The combined organic phases was dried over Na₂SO₄, filtered and concentrated *in vacuo*. The residue was dissolved in CH₂Cl₂ (15 mL) and trifluoroacetic acid (1.5 mL) was added. The reaction was stirred overnight at room temperature then concentrated *in vacuo*. The residue was purified by reverse phase HPLC (gradient 15-40% CH₃CN in H₂O containing 0.1% trifluoroacetic acid) to give COMPOUND 46 (213 mg, 76% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; 1 H NMR (400 MHz, CD₃OD) δ 0.92 (t, J = 7.32 Hz, 6H), 1.08-1.16 (m, 3H), 1.19-1.28 (m, 3H), 1.45-1.70 (m, 4H), 2.15-2.25 (m, 1H), 2.61 (dt, J = 15.72, 5.91 Hz, 4H), 3.22-3.34 (m, 6H), 3.48-3.58 (m, 2H), 6.90 (d, J =

7.62 Hz, 1H), 7.25-7.32 (m, 3H), 7.36 (d, J = 8.01 Hz, 3H), 7.56 (s, 1H). Found: C, 59.65; H, 5.46; N, 5.91. $C_{31}H_{35}N_3O_2$ x 1.8 CF_3CO_2H x 0.6 H_2O has C, 59.57; H, 5.49; N, 6.02%.

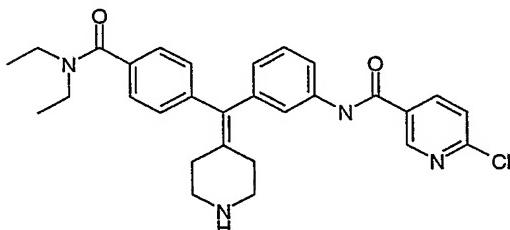
5 **COMPOUND 47:** *N*-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-1-methyl-1*H*-1,2,3-benzotriazole-5-carboxamide



Using the same method as for COMPOUND 46 and using INTERMEDIATE
10 6 (225 mg, 0.485 mmol) and 1-methyl-1*H*-1,2,3-benzotriazole-5-carbonyl chloride (142 mg, 0.728 mmol) afforded COMPOUND 47 (169 mg, 55% yield) as its TFA salt. This material was lyophilized from CH_3CN/H_2O to produce a colorless solid. Purity (HPLC): > 99%; 1H NMR (400 MHz, CD_3OD) δ 1.12 (br t, J = 6.59 Hz, 3H), 1.23 (br t, J = 7.03 Hz, 3H), 2.61 (t, J = 6.15 Hz, 2H), 2.67 (t, J = 5.96 Hz, 2H), 3.24-
15 3.34 (m, 6H), 3.53 (br q, J = 6.48 Hz, 2H), 4.38 (s, 3H), 6.95 (ddd, J = 7.62, 1.56, 1.07 Hz, 1H), 7.30 (d, J = 8.40 Hz, 2H), 7.33-7.39 (m, 3H), 7.54 (ddd, J = 8.15, 2.15, 1.03 Hz, 1H), 7.73 (t, J = 1.81 Hz, 1H), 7.88 (dd, J = 8.79, 0.78 Hz, 1H), 8.11 (dd, J = 8.69, 1.56 Hz, 1H), 8.59 (dd, J = 1.42, 0.83 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. $C_{30}H_{35}N_3O_2$ x 1.1 CF_3CO_2H x 1.0 H_2O has C, 63.79; H, 6.14; N, 6.72 %.

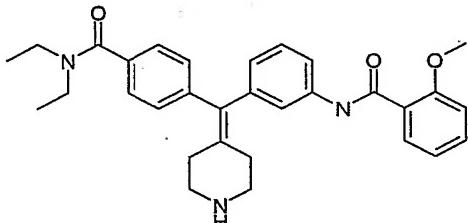
20

COMPOUND 48: 6-chloro-*N*-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-pyridinecarboxamide



Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 6-chloropyridine-3-carbonyl chloride (187 mg, 1.063 mol), except that the reaction was heated for 66 hours, afforded COMPOUND 48 (230 mg, 65% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.11 (br t, J = 5.86 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.60 (t, J = 5.86 Hz, 2H), 2.65 (t, J = 5.86 Hz, 2H), 3.23-3.33 (m, 6H), 3.48-3.58 (m, 2H), 6.96 (d, J = 7.62 Hz, 1H), 7.28 (d, J = 8.01 Hz, 2H), 7.32-7.40 (m, 3H), 7.50 (d, J = 8.01 Hz, 1H), 7.59 (d, J = 8.40 Hz, 1H), 7.68 (s, 1H), 8.27 (dd, J = 8.40, 2.54 Hz, 1H), 8.87 (d, J = 2.15 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. C₃₀H₃₅N₃O₂ x 1.1 CF₃CO₂H x 1.0 H₂O has C, 63.79; H, 6.14; N, 6.72 %.

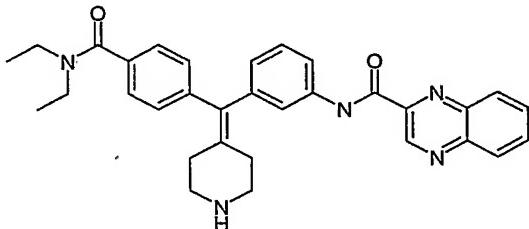
COMPOUND 49: N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-methoxy-benzamide



15

Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 2-methoxybenzoyl chloride (124 mg, 0.728 mmol) afforded COMPOUND 49 (221 mg, 75% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.61 (t, J = 6.05 Hz, 2H), 2.65 (t, J = 6.05 Hz, 2H), 3.23-3.34 (m, 6H), 3.53 (br q, J = 6.77 Hz, 2H), 4.00 (s, 3H), 6.92 (dt, J = 7.57, 1.29 Hz, 1H), 7.08 (dt, J = 7.42, 0.78 Hz, 1H), 7.18 (d, J = 8.40 Hz, 1H), 7.28-7.43 (m, 6H), 7.53 (ddd, J = 8.59, 7.23, 1.76 Hz, 1H), 7.76 (t, J = 1.76 Hz, 1H), 7.85 (dd, J = 7.81, 1.76 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. C₃₀H₃₅N₃O₂ x 1.1 CF₃CO₂H x 1.0 H₂O has C, 63.79; H, 6.14; N, 6.72 %.

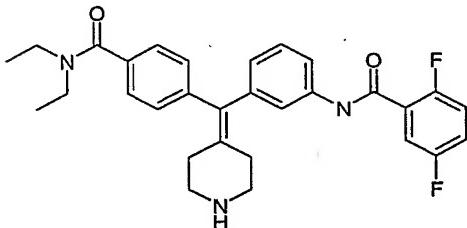
COMPOUND 50: *N*-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-quinoxalinecarboxamide



Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 2-quinoxaloyl chloride (140 mg, 0.728 mmol) afforded COMPOUND 50 (241 mg, 58% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.61 (t, J = 6.05 Hz, 2H), 2.65 (t, J = 6.05 Hz, 2H), 3.23-3.34 (m, 6H), 3.53 (br q, J = 6.77 Hz, 2H), 4.00 (s, 3H), 6.92 (dt, J = 7.57, 1.29 Hz, 1H), 7.08 (dt, J = 7.42, 0.78 Hz, 1H), 7.18 (d, J = 8.40 Hz, 1H), 7.28-7.43 (m, 6H), 7.53 (ddd, J = 8.59, 7.23, 1.76 Hz, 1H), 7.76 (t, J = 1.76 Hz, 1H), 7.85 (dd, J = 7.81, 1.76 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. C₃₀H₃₅N₃O₂ x 1.1 CF₃CO₂H x 1.0 H₂O has C, 63.79; H, 6.14; N, 6.72 %.

15

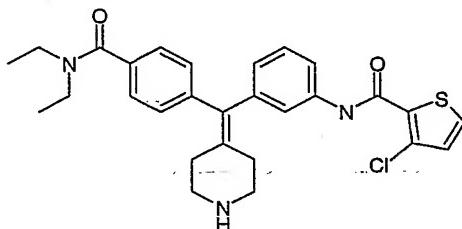
COMPOUND 51: *N*-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2,5-difluoro-benzamide



Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 2,5-difluorobenzoyl chloride (90 µL, 0.728 mmol) afforded COMPOUND 51 (233 mg, 78% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.23 (br t, J = 6.25 Hz,

3H), 2.61 (t, J = 6.05 Hz, 2H), 2.65 (t, J = 6.05 Hz, 2H), 3.23-3.34 (m, 6H), 3.53 (br q, J = 6.77 Hz, 2H), 4.00 (s, 3H), 6.92 (dt, J = 7.57, 1.29 Hz, 1H), 7.08 (dt, J = 7.42, 0.78 Hz, 1H), 7.18 (d, J = 8.40 Hz, 1H), 7.28-7.43 (m, 6H), 7.53 (ddd, J = 8.59, 7.23, 1.76 Hz, 1H), 7.76 (t, J = 1.76 Hz, 1H), 7.85 (dd, J = 7.81, 1.76 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. $C_{30}H_{35}N_3O_2 \times 1.1 CF_3CO_2H \times 1.0 H_2O$ has C, 63.79; H, 6.14; N, 6.72 %.

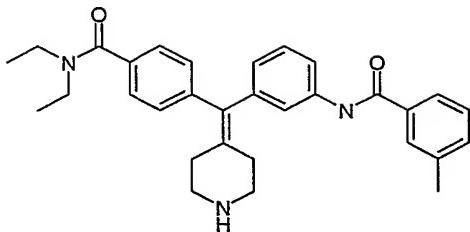
COMPOUND 52: 3-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-thiophenecarboxamide



10

Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 3-chlorothiophene-2-carbonyl chloride (132 mg, 0.728 mmol) afforded COMPOUND 52 (253 mg, 84% yield) as its TFA salt. This material was lyophilized from CH_3CN/H_2O to produce a colorless solid. Purity (HPLC): > 99%; 1H NMR (400 MHz, CD_3OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.61 (t, J = 6.05 Hz, 2H), 2.65 (t, J = 6.05 Hz, 2H), 3.23-3.34 (m, 6H), 3.53 (br q, J = 6.77 Hz, 2H), 4.00 (s, 3H), 6.92 (dt, J = 7.57, 1.29 Hz, 1H), 7.08 (dt, J = 7.42, 0.78 Hz, 1H), 7.18 (d, J = 8.40 Hz, 1H), 7.28-7.43 (m, 6H), 7.53 (ddd, J = 8.59, 7.23, 1.76 Hz, 1H), 7.76 (t, J = 1.76 Hz, 1H), 7.85 (dd, J = 7.81, 1.76 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. $C_{30}H_{35}N_3O_2 \times 1.1 CF_3CO_2H \times 1.0 H_2O$ has C, 63.79; H, 6.14; N, 6.72 %.

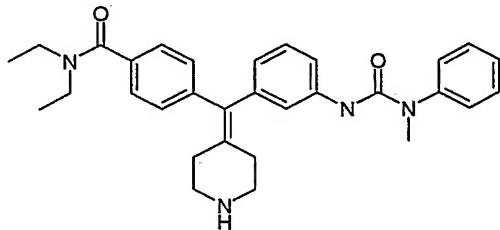
COMPOUND 53: N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-methyl-benzamide



Using the same method as for COMPOUND 46 and using INTERMEDIATE 6 (225 mg, 0.485 mmol) and 3-methylbenzoyl chloride (96 μ L, 0.728 mmol) afforded 5 COMPOUND 53 (227 mg, 79% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.12 (br t, J = 6.64 Hz, 3H), 1.23 (br t, J = 6.25 Hz, 3H), 2.61 (t, J = 6.05 Hz, 2H), 2.65 (t, J = 6.05 Hz, 2H), 3.23-3.34 (m, 6H), 3.53 (br q, J = 6.77 Hz, 2H), 4.00 (s, 3H), 6.92 (dt, J = 7.57, 1.29 Hz, 1H), 7.08 (dt, J = 7.42, 0.78 Hz, 1H), 10 7.18 (d, J = 8.40 Hz, 1H), 7.28-7.43 (m, 6H), 7.53 (ddd, J = 8.59, 7.23, 1.76 Hz, 1H), 7.76 (t, J = 1.76 Hz, 1H), 7.85 (dd, J = 7.81, 1.76 Hz, 1H). Found: C, 63.88; H, 6.13; N, 6.66. C₃₀H₃₅N₃O₂ x 1.1 CF₃CO₂H x 1.0 H₂O has C, 63.79; H, 6.14; N, 6.72 %.

COMPOUND 54: N,N-diethyl-4-[[3-

15 **[[(methylphenylamino)carbonyl]amino]phenyl]-4-piperidinylidenemethyl]-benzamide**

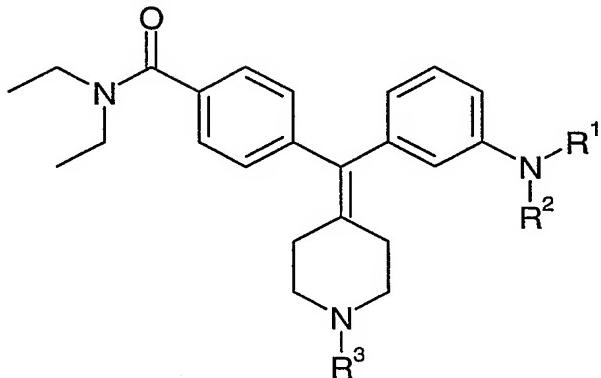


To a solution of INTERMEDIATE 6 (225 mg, 0.485 mmol) and N-methyl-N-phenylcarbamoyl chloride (370 mg, 2.183 mmol) in 1,2-dichloroethane (20 mL) was added triethylamine (0.203 mL, 1.456 mmol). The reaction was heated for 6 days at 20 50 °C, diluted with CH₂Cl₂ and washed with saturated aqueous NaHCO₃ (1x). The layers were separated and the aqueous layer was extracted with additional CH₂Cl₂ (1x). The combined organic phases was dried over Na₂SO₄, filtered and concentrated

in vacuo. The residue was dissolved in CH₂Cl₂ (15 mL) and trifluoroacetic acid (1.5 mL) was added. The reaction was stirred overnight at room temperature then concentrated *in vacuo*. The residue was purified by reverse phase HPLC (gradient 10-40% CH₃CN in H₂O containing 0.1% trifluoroacetic acid) to give COMPOUND 54 (213 mg, 76% yield) as its TFA salt. This material was lyophilized from CH₃CN/H₂O to produce a colorless solid. Purity (HPLC): > 99%; ¹H NMR (400 MHz, CD₃OD) δ 1.11 (br t, J = 6.25 Hz, 3H), 1.23 (br t, J = 6.74 Hz, 3H), 2.58 (dt, J = 12.25, 6.08 Hz, 4H), 3.19-3.28 (m, 6H), 3.30 (s, 3H), 3.53 (br q, J = 7.16 Hz, 2H), 6.79 (dt, J = 7.66, 1.42 Hz, 1H), 7.12 (ddd, J = 8.20, 2.15, 1.17 Hz, 1H), 7.19 (d, J = 7.62 Hz, 1H), 7.22-7.26 (m, 2H), 7.29 (t, J = 1.76 Hz, 1H), 7.31-7.36 (m, 5H), 7.42-7.49 (m, 2H). Found: C, 59.65; H, 5.46; N, 5.91. C₃₁H₃₅N₃O₂ x 1.8 CF₃CO₂H x 0.6 H₂O has C, 59.57; H, 5.49; N, 6.02%.

What is claimed is :

1. A compound of formula I, or a pharmaceutically acceptable salt thereof:



5

I

wherein

- R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₉heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl,
 C₂₋₉heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, R⁸-C(=O)-,
 10 R⁸-S(=O)₂-, R⁸-S(=O)-, R⁸-NHC(=O)-, R⁸-C(=S)- and R⁸-NH-C(=S)-, wherein R⁸ is
 selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₉heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₉heteroaryl-
 C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl,
 C₆₋₁₀aryl, C₂₋₉heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₉heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl,
 and C₃₋₁₀cycloalkyl-C₁₋₄alkyl used in defining R¹ and R⁸ are optionally substituted
 15 with one or more groups selected from -R, -NO₂, -OR, -Cl, -Br, -I, -F, -CF₃,
 -C(=O)R, -C(=O)OH, -NH₂, -SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(=O)R, -CN,
 -OH, -C(=O)OR, -C(=O)NR₂, -NRC(=O)R, and -NRC(=O)-OR, wherein R is,
 independently, selected from -H, C₁₋₆alkyl and phenyl;

- R² is selected from -H and C₁₋₆alkyl optionally substituted with one or more
 20 groups selected from halogen, -CF₃, -OH, C₁₋₃alkoxy, and halogen, or R¹ and R² are
 C₁₋₃alkylene that together form a portion of a ring; and

- R³ is selected from -H, C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and
 C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl,
 and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups
 25 selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

2. A compound according to claim 1, wherein

R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

R² is selected from -H and C₁₋₃alkyl; and

10 R³ is selected from -H and C₁₋₆alkyl-O-C(=O)-.

3. A compound according to claim 2,

wherein R¹ is R⁹-CH₂-, wherein R⁹ is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, 15 pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy and halogen; and

R² and R³ are hydrogen.

20

4. A compound according to claim 3,

wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl, optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen.

25

5. A compound according to claim 4, wherein

wherein R⁹ is selected from benzyl, phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl and thiazolyl.

30 6. A compound according to claim 1, wherein

R¹ is selected from C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally

substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

R² is -H or C₁₋₃alkyl; and

R³ is -H, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said
5 C₁₋₆alkyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one
or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and
halogen.

7. A compound according to claim 6, wherein

10 R¹ is selected from 1-propyl, 2-propyl, 1-butyl, 2-butyl, t-butyl, 2-methyl-1-
propyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, and cyclononyl;
R² is selected from -H, methyl, ethyl, 1-propyl and 2-propyl; and
R³ is selected from -H, methyl, ethyl, allyl, 3,3-dimethyl-allyl,
cyclopropylmethyl, 2-methoxy-ethyl, and 3-methoxy-1-propyl.

15 8. A compound according to claim 1, wherein

R¹ is selected from R⁸-C(=O)-, R⁸-S(=O)₂-, R⁸-S(=O)-, R⁸-NHC(=O)-, R⁸-
C(=S)- and R⁸-NH-C(=S)-, wherein R⁸ is selected from C₃₋₆alkyl, C₆₋₁₀aryl,
C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₂₀
cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄
alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are
optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and
halogen;

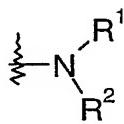
R² is -H; and

25 R³ is selected from -H and C₁₋₆alkyl-O-C(=O)-.

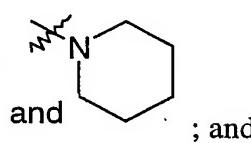
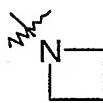
9. A compound according to claim 8, wherein

R⁸ is selected from phenyl, benzyl, phenethyl and cyclohexyl, wherein said
phenyl, benzyl, phenethyl and cyclohexyl are optionally substituted with one or more
30 groups selected from methyl, methoxy and halogen.

10. A compound according to claim 1, wherein



of formula I is selected from



and ; and
 R^3 is selected from -H and C_{1-6} alkyl-O-C(=O)-.

5 11. A compound selected from:

- 1) 4-[[3-(benzylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 2) N,N-diethyl-4-[[3-[(3-furylmethyl)amino]phenyl](piperidin-4-ylidene)methyl]benzamide,
- 3) N,N-diethyl-4-(piperidin-4-ylidene){3-[(thien-3-ylmethyl)amino]phenyl}methyl)benzamide,
- 4) N,N-diethyl-4-[[3-[(2-phenylethyl)amino]phenyl](piperidin-4-ylidene)methyl]benzamide,
- 5) 4-[[3-[(4-chlorobenzyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 6) N,N-diethyl-4-[piperidin-4-ylidene(3-{{[3-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,
- 7) 4-[[3-[(2-chlorobenzyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 8) N,N-diethyl-4-[piperidin-4-ylidene(3-{{[4-(trifluoromethyl)benzyl]amino}phenyl)methyl]benzamide,
- 9) N,N-diethyl-4-[[3-[(2-furylmethyl)amino]phenyl](piperidin-4-ylidene)methyl]benzamide,
- 10) N,N-diethyl-4-(piperidin-4-ylidene){3-[(thien-2-ylmethyl)amino]phenyl}methyl)benzamide,
- 11) 4-[[3-[(cyclohexylmethyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 12) N,N-diethyl-4-{piperidin-4-ylidene[3-(propylamino)phenyl]methyl}benzamide,

- 13) 4-[[3-(cyclohexylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 14) 4-[[3-(cyclopentylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 5 15) 4-[[3-(cycloheptylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 16) 4-[[3-[cyclopentyl(methyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 17) 4-[[3-(benzoylamino)phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 10 18) N,N-diethyl-4-[[3-[(phenylacetyl)amino]phenyl](piperidin-4-ylidene)methyl]benzamide,
- 19) 4-[[3-[(cyclohexylcarbonyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 15 20) 4-[[3-[(cyclohexylacetyl)amino]phenyl](piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 21) 4-[(3-{{(2-chlorophenyl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 22) 4-[(3-{{(3-chlorophenyl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 20 23) N,N-diethyl-4-[(3-{{(5-methylthien-2-yl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 24) 4-[(3-{{(5-chlorothien-2-yl)acetyl}amino}phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 25 25) N,N-diethyl-4-[(3-{{(2S)-2-phenylpropanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 26) N,N-diethyl-4-[(3-{{(2R)-2-phenylpropanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 27) N,N-diethyl-4-[(3-{{(2S)-2-phenylbutanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 30 28) N,N-diethyl-4-[(3-{{(2R)-2-phenylbutanoyl}amino}phenyl)(piperidin-4-ylidene)methyl]benzamide,

- 29) 4-[{3-[benzoyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 30) 4-[{3-[(anilinocarbonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 5 31) 4-[{(benzylamino)carbonyl}amino]phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 32) N-{3-[{4-[(diethylamino)carbonyl]phenyl}(piperidin-4-ylidene)methyl]phenyl}piperidine-1-carboxamide,
- 33) N,N-diethyl-4-[{3-[(phenylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 10 34) 4-[{3-[(benzylsulfonyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 35) 4-[(3-anilinophenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 36) N,N-diethyl-4-[{3-[methyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 15 37) N,N-diethyl-4-[{3-[ethyl(phenyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 38) N,N-diethyl-4-[{(1S)-1-phenylethyl}amino]phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 20 39) N,N-diethyl-4-[{(1R)-1-phenylethyl}amino]phenyl)(piperidin-4-ylidene)methyl]benzamide,
- 40) 4-[{(1R)-1-cyclohexylethyl}amino]phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 41) 4-[{(1S)-1-cyclohexylethyl}amino]phenyl)(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 25 42) N,N-diethyl-4-[{3-[(1-methyl-1-phenylethyl)amino]phenyl}(piperidin-4-ylidene)methyl]benzamide,
- 43) 4-[{3-[cyclohexyl(methyl)amino]phenyl}(piperidin-4-ylidene)methyl]-N,N-diethylbenzamide,
- 30 44) N,N-diethyl-4-[piperidin-4-ylidene(3-piperidin-1-ylphenyl)methyl]benzamide,
- 45) N,N-diethyl-4-[piperidin-4-ylidene(3-pyrrolidin-1-ylphenyl)methyl]benzamide,

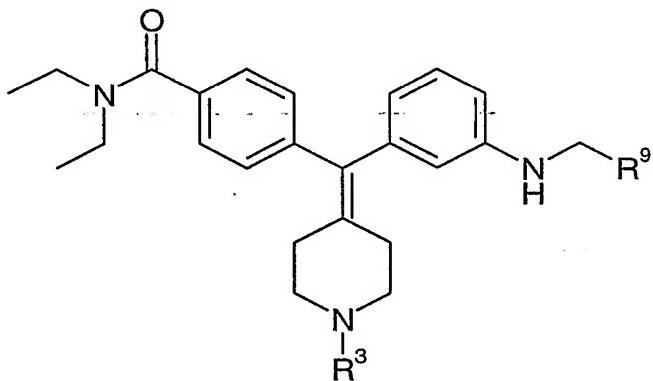
- 46) N,N-diethyl-4-[[3-[(2-ethyl-1-oxobutyl)amino]phenyl]-4-piperidinylidenemethyl]-benzamide,
- 47) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-1-methyl-1H-1,2,3-benzotriazole-5-carboxamide,
- 5 48) 6-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-pyridinecarboxamide,
- 49) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-methoxy-benzamide,
- 10 50) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-quinoxalinecarboxamide,
- 51) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2,5-difluoro-benzamide,
- 52) 3-chloro-N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-2-thiophenecarboxamide,
- 15 53) N-[3-[[4-[(diethylamino)carbonyl]phenyl]-4-piperidinylidenemethyl]phenyl]-3-methyl-benzamide,
- 54) N,N-diethyl-4-[[3-[(methylphenylamino)carbonyl]amino]phenyl]-4-piperidinylidenemethyl]-benzamide, and pharmaceutically acceptable salts thereof.
- 20 12. A compound according to any one of claims 1-11 for use as a medicament.
13. The use of a compound according to any one of claims 1-11 in the manufacture of a medicament for the therapy of pain, anxiety or functional gastrointestinal disorders.
- 25 14. A pharmaceutical composition comprising a compound according to any one of claims 1-11 and a pharmaceutically acceptable carrier.
15. A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-11.

16. A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-11.

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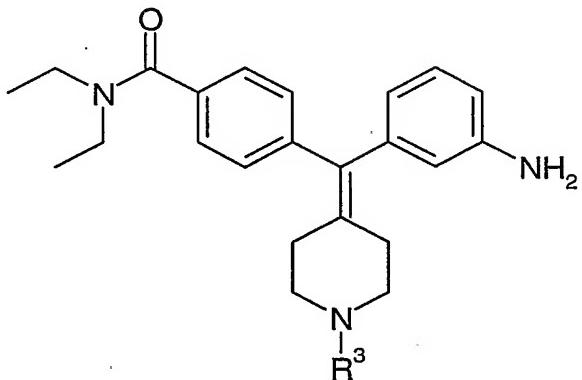
17. A method for the therapy of anxiety in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-11.

10 18. A process for preparing a compound of formula III,

III

comprising:

reacting a compound of formula II,



15

II

with R⁹-CHO in the presence of a reducing agent to form the compound of formula III,

wherein

86

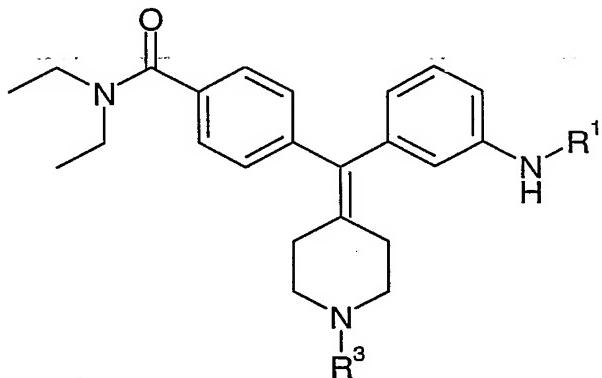
R^9 is selected from phenyl, pyridyl, thienyl, furyl, imidazolyl, triazolyl, pyrrolyl, thiazolyl, N-oxido-pyridyl, benzyl, pyridylmethyl, thienylmethyl, furylmethyl, imidazolylmethyl, triazolylmethyl, pyrrolylmethyl, thiazolylmethyl and N-oxido-pyridylmethyl, optionally substituted with one or more groups selected from

5 C_{1-4} alkyl, halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, phenoxy and halogen; and

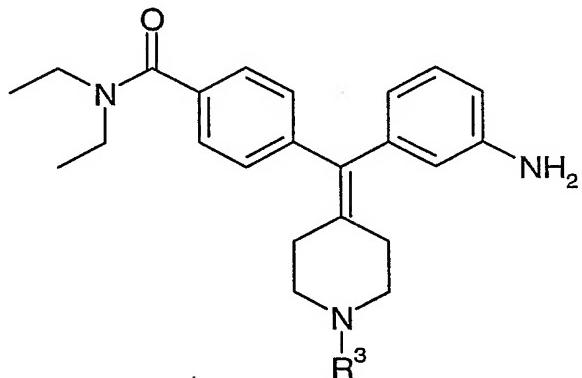
R^3 is selected from C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy and halogen.

10

19. A process for preparing a compound of formula IV,

IV

comprising: reacting a compound of formula II,



15

II

with R^1-X to form the compound of formula IV,

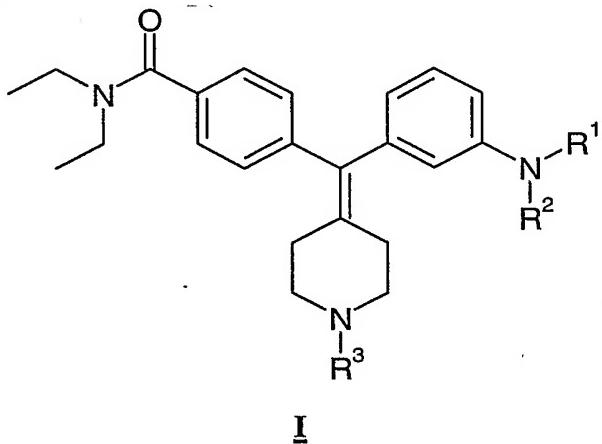
wherein

X is halogen;

R^1 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl, wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, C_{3-10} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-4} alkyl, halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, phenoxy, and halogen; and

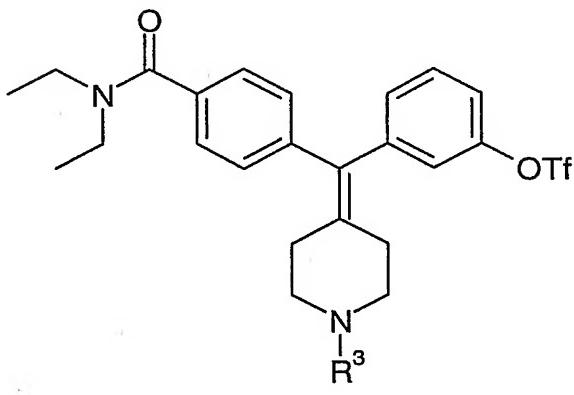
R^3 is selected from C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl-O-C(=O)-, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy and halogen.

20. A process for preparing a compound of formula I,



I

15 comprising: reacting a compound of formula V,



V

with R^1R^2NH to form the compound of formula I,

wherein

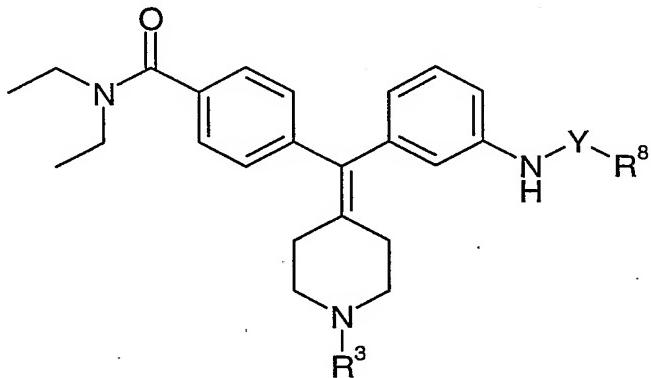
88

R¹ is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl, wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen;

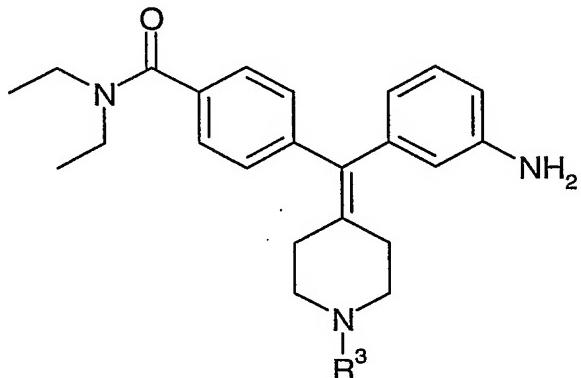
5 R² is selected from -H and C₁₋₆alkyl optionally substituted with one or more groups selected from halogen, -CF₃, -OH, C₁₋₃alkoxy, and halogen, or R¹ and R² are C₁₋₃alkylene that together form a portion of a ring; and

10 R³ is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

15 21. A process for preparing a compound of formula VI,

VI

comprising: reacting a compound of formula VII,



89**VII**

with R^8 -Y-X or R^8 -Y-O-Y-R⁸ to form the compound of formula VI:

wherein

X is halogen;

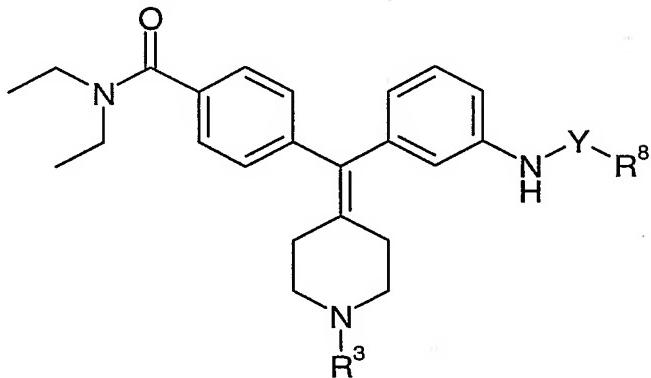
5 Y is selected from -C(=O)- and -S(=O)₂-;

R^8 is selected from C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl; wherein said C₃₋₆alkyl, C₆₋₁₀aryl, C₂₋₆heteroaryl, C₆₋₁₀aryl-C₁₋₄alkyl, C₂₋₆heteroaryl-C₁₋₄alkyl, C₃₋₁₀cycloalkyl, and C₃₋₁₀cycloalkyl-C₁₋₄alkyl are optionally substituted with C₁₋₄alkyl, halogen, -CF₃, -OH, C₁₋₃alkoxy, phenoxy, and halogen; and

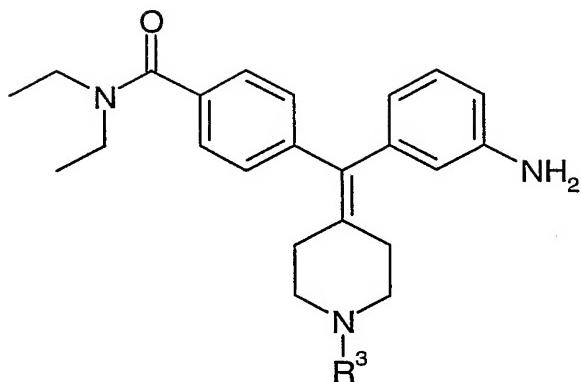
10 R^3 is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

15

22. A process for preparing a compound of formula VIII,

**VIII**

comprising: reacting a compound of formula VII,

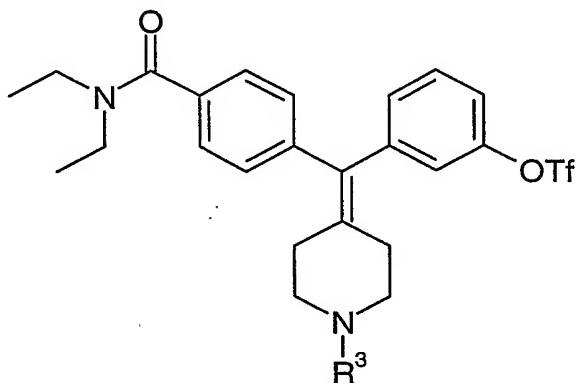
90**VII**

with R^8 -Z to form the compound of formula VIII:

wherein

- 5 Z is selected from -NCO and -NCS;
- Y is selected from $-C(=O)NH-$ and $-C(=S)NH-$;
- 10 R^8 is selected from C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl; wherein said C_{3-6} alkyl, C_{6-10} aryl, C_{2-6} heteroaryl, C_{6-10} aryl- C_{1-4} alkyl, C_{2-6} heteroaryl- C_{1-4} alkyl, C_{3-10} cycloalkyl, and C_{3-10} cycloalkyl- C_{1-4} alkyl are optionally substituted with C_{1-4} alkyl, halogen, $-CF_3$, $-OH$, C_{1-3} alkoxy, phenoxy, and halogen; and
- 15 R^3 is selected from C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl, wherein said C_{1-6} alkyl- $O-C(=O)-$, C_{1-6} alkyl, C_{3-6} cycloalkyl, and C_{3-6} cycloalkyl- C_{1-4} alkyl are optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy and halogen.

23. A compound of formula V,

**V**

wherein

- R³ is selected from C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl, wherein said C₁₋₆alkyl-O-C(=O)-, C₁₋₆alkyl, C₃₋₆cycloalkyl, and C₃₋₆cycloalkyl-C₁₋₄alkyl are optionally substituted with one or more groups
5 selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy and halogen.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2003/001705

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 211/70, A61K 31/4465, A61P 25/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CHEM. ABS DATA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9828275 A1 (ASTRA PHARMA INC.), 2 July 1998 (02.07.1998) ---	1-22

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

20 February 2004

Date of mailing of the international search report

24 -02- 2004

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INTERNATIONAL SEARCH REPORT
Information on patent family members

24/12/2003

International application No.
PCT/SE 2003/001705

WO	9828275	A1	02/07/1998	AU	737999	B	06/09/2001
				AU	5351298	A	17/07/1998
				BR	9714055	A	09/05/2000
				CN	1246111	A	01/03/2000
				CZ	9902199	A	17/11/1999
				EE	3824	B	15/08/2002
				EE	9900256	A	15/12/1999
				EP	0946511	A	06/10/1999
				ID	22074	A	00/00/0000
				IL	130535	D	00/00/0000
				JP	2001507350	T	05/06/2001
				NO	313670	B	11/11/2002
				NO	993022	A	20/08/1999
				NZ	336029	A	30/03/2001
				PL	334374	A	28/02/2000
				RU	2193029	C	20/11/2002
				SE	9604785	D	00/00/0000
				SK	76299	A	08/11/1999
				SK	283211	B	04/03/2003
				TR	9901417	T	00/00/0000
				TW	548271	B	00/00/0000
				US	6187792	B	13/02/2001
				US	6455545	B	24/09/2002
				US	2001021715	A	13/09/2001
				US	2003149023	A	07/08/2003
				ZA	9711059	A	22/06/1998
				HU	0000610	A	28/09/2000
				SE	9702535	D	00/00/0000

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/SE 2003/001705**Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: **15-17**
because they relate to subject matter not required to be searched by this Authority, namely:
see next sheet
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 2003/001705

Box No. IV Text of the abstract (Continuation of item 5 of the first sheet)

Claims 15-17 relate to methods of treatment of the human or animal body by surgery or by therapy or diagnostic methods practised on the human or animal body (PCT Rule 39.1(iv)). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds or compositions.